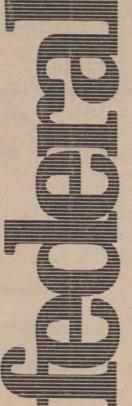
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Tuesday September 1, 1987

Briefings on How To Use the Federal Register— For information on briefings in Washington, DC, see announcement on the inside cover of this issue.



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To provide the public with access to information WHY: necessary to research Federal agency regulations which directly affect them. There will be no discussion of specific agency regulations.

WASHINGTON, DC

September 29, at 9 a.m. WHEN: WHERE: Office of the Federal Register, First Floor Conference Room,

1100 L Street NW., Washington, DC. RESERVATIONS: Janice Booker, 202-523-5239

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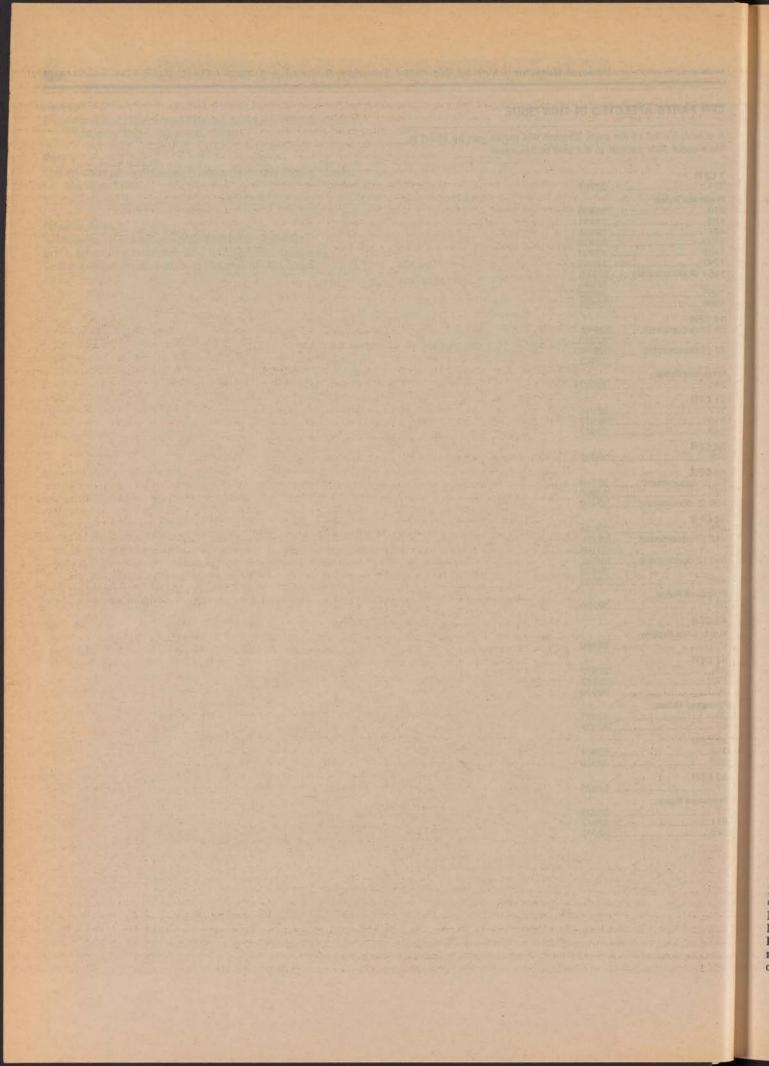
Reader Aids

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Tuesday, September 1, 1987

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The Code of Federal Regulations is sold by the Superintendent of Documents. Prices of new books are listed in the first FEDERAL REGISTER issue of each

week.

DEPARTMENT OF AGRICULTURE

Animal and Plant Health Inspection Service

7 CFR Part 301

[Docket No. 87-077]

Imported Fire Ant Regulated Areas

AGENCY: Animal and Plant Health Inspection Service, USDA.

ACTION: Interim rule.

SUMMARY: We are amending the list of generally infested areas under the imported fire ant quarantine and regulations by adding areas in 6 counties in Alabama, 2 counties in Arkansas, 11 counties in Georgia, 2 counties in Mississippi, 11 counties in North Carolina, and 42 counties in Texas.

We are also amending the imported fire ant quarantine and regulations by quarantining the state of Oklahoma, and by designating areas in three counties in Oklahoma as generally infested areas.

In addition, we are making nonsubstantive, editorial changes.

This action imposes certain restrictions on the interstate movement of regulated articles and expands the regulated area. It is necessary to prevent the artificial spread of the imported fire

DATE: Interim rule effective September 1, 1987. Consideration will be given only to comments postmarked or received on or before November 2, 1987.

ADDRESSES: Send an original and two copies of written comments to Steven B. Farbman, Assistant Director, Regulatory Coordination, APHIS, USDA, Room 728, Federal Building, Hyattsville, MD 20782. Please state that your comments refer to Docket Number 87-077. Comments received may be inspected at Room 728 of the Federal Building between 8 a.m.

and 4:30 p.m., Monday through Friday, except holidays.

FOR FURTHER INFORMATION CONTACT: Charles H. Bare, Staff Officer, Field Operations Support Staff, PPQ, APHIS, USDA, Room 663, Federal Building. Hyattsville, MD 20782, 301-436-8295.

SUPPLEMENTARY INFORMATION:

Background

The imported fire ant quarantine and regulations contained in 7 CFR 301.81, et. seq., and (referred to below as the regulations) restrict the interstate movement of regulated articles from regulated areas in designated states to prevent the artificial spread of the imported fire ant. The improted fire ant (Solenopsis spp.) is an insect that interferes with farming operations, can cause damage to certain crops, and is a pest of livestock, pets, and people in rural and urban areas. The quarantined states are: Alabama, Arkansas, Florida, Georgia, Louisiana, Mississippi, North Carolina, Puerto Rico, South Carolina, and Texas.

Under the regulations, an area is designated as a regulated area if the imported fire ant has been found there. or if reason exists to believe the imported fire ant is present there.

Regulated areas are designated as either generally infested areas or suppressive areas. Suppressive areas are those areas where eradication of the imported fire ant is being undertaken as an objective. Generally infested areas are all other regulated areas.

Restrictions are imposed on the interstate movement of regulated articles from regulated areas to prevent the artificial movement of imported fire ant into noninfested areas, and to prevent further infestation of suppressive areas.

Designation of Areas as Generally Infested Areas

We are amending § 301.81(a) of the regulations by adding Oklahoma to the list of quarantined states. We are also amending § 301.81-2a by designating portions of 3 Oklahoma counties as generally infested areas.

We are also amending the list of generally infested areas in § 301.81-2a of the regulations by adding areas in quarantined states as follows: Cherokee, De Kalb, Jackson, Madison, Marshall, and Morgan Counties in Alabama: Dallas and Howard Counties in

Arkansas; Banks, Barrow, Clarke, Elbert, Gordon, Jackson, Lincoln, Madison, Oglethorpe, Whitfield, and Wilkes Counties in Georgia; Marshall and Tippah Counties in Mississippi; Beaufort, Bladen, Craven, Duplin, Hyde. Lenoir, Richmond, Robeson, Sampson, Scotland, and Union Counties in North Carolina; and Bandera, Bee, Bosque, Bowie, Burnet, Camp, Cass, Comanche, Cooke, Corvell, Duval, Eastland, Erath. Fannin, Franklin, Frio, Grayson, Henderson, Hood, Hunt, Jim Wells, Karnes, Kaufman, Kimble, Lampasas, Llano, Medina, Morris, Parker, Rains, Somervell, Taylor, Titus, Tom Green, Uvalde, Van Zandt, Wharton, Wichita, Williamson, Wilson, Wise, and Young Counties in Texas.

See the rule portion of this document for specific descriptions of newly infested areas.

These actions are necessary because surveys conducted by inspectors of the United States Department of Agriculture and officials of state agencies establish that the imported fire ant has spread to these areas. Eradication of the imported fire ant is not being undertaken as an objective in these areas, and thereforeas an emergency measure—we are adding them to the list of imported fire ant generally infested areas.

Emergency Action

William F. Helms, Deputy Administrator of the Animal and Plant Health Inspection Service for Plant Protection and Quarantine, has determined that an emergency situation exists which warrants publication of this interim rule without prior opportunity for public comment. Because the imported fire ant could be spread artificially to noninfested areas of the United States, it is necessary to act immediately to control its spread.

Since prior notice and other public procedures with respect to this interim rule are impracticable and contrary to the public interest under these emergency conditions, there is good cause under 5 U.S.C. 553(d)(3) for making this interim rule effective less than 30 days after publication of this document in the Federal Register. We will consider comments postmarked or received within 60 days of publication of this interim rule in the Federal Register. Any amendments we make to this interim rule as a result of these comments will be published in the

Federal Register as soon as possible following the close of the comment period.

Executive Order 12291 and Regulatory Flexibility Act

We are issuing this rule in conformance with Executive Order 12291, and we have determined that it is not a "major rule." Based on information compiled by the Department, we have determined that this rule will have an estimated annual effect on the economy of approximately \$37,500; will not cause a major increase in costs or prices for consumers, individual industries, Federal, State, or local government agencies, or geographic regions; and will not cause a significant adverse effect on competition, employment, investment, productivity, innovation, or on the ability of United States-based enterprises to compete with foreignbased enterprises in domestic or export markets.

For this action, the Office of Management and Budget has waived the review process required by Executive Order 12291.

This action affects the interstate movement of regulated articles from specified areas in Alabama, Arkansas, Georgia, Mississippi, North Carolina, Oklahoma, and Texas. Thousands of small entities move these articles interstate from the above mentioned States, and many more thousands of small entities move these articles interstate from other States.

However, based on information compiled by the Department, we have determined that approximately 896 small entities move these articles interstate from the specified areas in those States. Further, the overall economic impact from this action is estimated to be approximately \$37,500.

Under these circumstances, the Administrator of the Animal and Plant Health Inspection Service has determined that this action will not have a significant economic impact on a substantial number of small entities.

Executive Order 12372

This program/activity is listed in the Catalog of Federal Domestic Assistance under No. 10.025 and is subject to the provisions of Executive Order 12372, which requires intergovernmental consultation with state and local officials. (See 7 CFR Part 3015, Subpart V.)

Paperwork Reduction Act

This interim rule contains no information collection or recordkeeping requirements under the Paperwork

Reduction Act of 1980 (44 U.S.C. 3501 et seq.).

List of Subjects in 7 CFR Part 301

Agricultural commodities, Imported fire ant, Plant diseases, Plant pests, Plants (Agriculture), Quarantine, Transportation.

Accordingly, 7 CFR Part 301 is amended as follows:

PART 301—DOMESTIC QUARANTINE NOTICES

1. The authority citation for Part 301 continues to read as follows:

Authority: 7 U.S.C. 150dd, 150ee, 150ff, 161, 162, and 164–167; 7 CFR 2.17, 2.51, and 371.2(c).

§ 301.81 [Amended]

- 2. Section 301.81, paragraph (a), is amended by adding "Oklahoma" immediately before "Puerto Rico."
- 3. Section 301.81-2a is revised to read as follows:

§ 301.81-2a Regulated areas; suppressive and generally infested areas.

The civil divisions and parts of civil divisions described below are designated as imported fire ant regulated areas within the meaning of the provisions of this subpart; and these regulated areas are hereby divided into generally infested areas or suppressive areas as indicated below:

Alabama

(1) Generally infested areas. Autauga County. The entire county. Baldwin County. The entire county. Barbour County. The entire county. Bibb County. The entire county. Blount County. The entire county. Bullock County. The entire county. Butler County. The entire county. Calhoun County. The entire county. Chambers County. The entire county. Cherokee County. The entire county. Chilton County. The entire county. Choctaw County. The entire county. Clarke County. The entire county. Clay County. The entire county. Cleburne County. The entire county. Coffee County. The entire county. Colbert County. The entire county. Conecuh County. The entire county. Coosa County. The entire county. Covington County. The entire county. Crenshaw County. The entire county. Cullman County. The entire county.

Dale County. The entire county. Dallas County. The entire county. De Kalb County. T. 8 and 9 S., R. 5 E.; W. 1/ 2 T. 8 and 9 S., R. 6 E.; secs. 21, 22, 23, 24, 25, 26, 27, 28, 33, 34, 35, and 36, T. 9 S., R. 6 E.; T. 8 and 9 S., R. 7 E.; T. 8 and 9 S., R. 8 E.; T. 7 and 8 S., R. 9 E.; T. 7 S., R. 10 E.; secs. 26 and 35, T. 7 S., R. 7 E.; secs. 9, 10, 15, and 16, T. 7 S., R. 8 E.

Elmore County. The entire county. Escambia County. The entire county.

Etowah County. The entire county.
Fayette County. The entire county.
Franklin County. The entire county.
Geneva County. The entire county.
Greene County. The entire county.
Hale County. The entire county.
Henry County. The entire county.
Houston County. The entire county.
Jackson County. Secs. 11, 12, 13, and 14, T.
5 S., R. 4 E.; secs. 7, 8, 17, and 18, T. 5 S.; R. 5
F.

Jefferson County. The entire county.

Lamar County. The entire county.

Lauderdale County. T. 2 S., R. 10, 11, 12, 13, 14, and 15 W.; T. 3 S., R. 9, 10, 11, 12, and 13 W.

Lawrence County. The entire county.

Lee County. The entire county.

Limestone County. S. ½ T. 3 S., R. 6 W.;

W. ½ T. 4 S., R. 5 W.; T. 4 S., R. 6 W.; T. 5 S.,

R 4 W.

Lowndes County. The entire county. Macon County. The entire county. Madison County. That portion of the county south of the north line of T. 5 S. Marengo County. The entire county. Marion County. The entire county. Marshall County. The entire county. Mobile County. The entire county. Monroe County. The entire county. Montgomery County. The entire county. Morgan County. The entire county. Perry County. The entire county. Pickens County. The entire county. Pike County. The entire county. Randolph County. The entire county. Russell County. The entire county. St. Clair County. The entire county Shelby County. The entire county. Sumter County. The entire county. Talladega County. The entire county. Tallapeosa County. The entire county. Tuscaloosa County. The entire county. Walker County. The entire county. Washington County. The entire county. Wilcox County. The entire county. Winston County. The entire county. (2) Suppressive areas. None.

Arkansas

(1) Generally infested areas.
Ashley County. The entire county.
Bradley County. The entire county.
Calhoun County. The entire county.
Chicot County. The entire county.
Cleveland County. The entire county.
Columbia County. The entire county.
Dallas County. The entire county.
Desha County. That portion of the county south of the south line of T. 10 S.
Drew County. The entire county.

Drew County. The entire county.

Hempstead County. That portion of the county south of Interstate 30 including all of the incorporated city limits of Hope.

Howard County. T. 9 and 10 S., R. 27 W. Jefferson County. Secs. 30, 31, and that part of secs. 29, 32, and 33 south and east of the Arkansas River of T. 5 S., R. 8 W.; secs. 25 and 36 of T. 5 S., R. 9 W.; secs. 8, 9, 10, 11, 14, 15, 16, and 17 of T. 6 S., R. 8 W.

Lafayette County. The entire county.
Lincoln County. That portion of the county south of State Road 114 and west of State
Road 81, including all of the incorporated city limits of Palmyra and Star City.

Little River County. The entire county.

Miller County. The entire county.

Nevada County. That portion of the county south of the south line of T. 10 S., and the Little Missouri River.

Quachita County. The entire county. Union County. The entire county. (2) Suppressive areas. None.

Florida

(1) Generally infested areas. The entire state.

(2) Suppressive areas. None.

Georgia

(1) Generally infested areas.
Appling County. The entire county.
Atkinson County. The entire county.
Bacon County. The entire county.
Baker County. The entire county.
Baldwin County. The entire county.
Banks County. That portion of the county
within Georgia Military Districts 208, 265, 448,
468, 912, 1206, 1210, and 1464.

Barrow County. The entire county.
Bartow County. The entire county.
Ben Hill County. The entire county.
Berrien County. The entire county.
Bibb County. The entire county.
Bleckley County. The entire county.
Brooks County. The entire county.
Brooks County. The entire county.
Bryan County. The entire county.
Bulloch County. The entire county.
Buthe County. The entire county.
Buths County. The entire county.
Calhoun County. The entire county.
Candler County. The entire county.
Carroll County. The entire county.
Charlton County. The entire county.
Charlton County. The entire county.
Charlton County. The entire county.
Chatham County. The entire county.
Chatham County. The entire county.
Chattachoochee County. The entire county.
Chattacoga County. That portion of the
county within Georgia Militia Districts 925,
961, 968, 1083, 1216, and 1484.

Cherokee County. That portion of the county within Georgia Militia District 817. Clarke County. The entire county. Clay County. The entire county. Clayton County. The entire county. Clinch County. The entire county. Cobb County. The entire county. Coffee County. The entire county. Colquitt County. The entire county. Columbia County. The entire county. Cook County. The entire county. Coweta County. The entire county. Crawford County. The entire county. Crisp County. The entire county. Decatur County. The entire county. De Kalb County. The entire county. Dodge County. The entire county. Dooly County. The entire county. Dougherty County. The entire county. Douglas County. The entire county. Early County. The entire county. Echols County. The entire county.

Effingham County. The entire county. Elbert County. That portion of the county within Georgia Militia Districts 190, 191, 192,

Emanuel County. The entire county.

Evans County. The entire county.

Fayette County. The entire county.

Floyd County. That portion of the county
within Georgia Militia Districts 829, 855, 859,

919, 923, 924, 962, 1048, 1059, 1120, 1453, 1478, 1504, 1562, 1688, 1719, and 1822.

Forsyth County. That portion of the county within Georgia Militia Districts 879, 1276, and 795.

Fulton County. The entire county.

Glascock County. The entire county.

Glynn County. The entire county.

Gordon County. That portion of the county within Georgia Militia Districts 849, 856, 973, 930, 1054, 1055, 1056, 1064, and 1595.

Grady County. The entire county.

Greene County. The entire county.

Gwinnett County. The entire county.

Hall County. That portion of the county within Georgia Militia Districts 413, 1270, and

Hancock County. The entire county. Haralson County. The entire county. Harris County. The entire county. Heard County. The entire county. Henry County. The entire county. Houston County. The entire county. Irwin County. The entire county. Jackson County. The entire county. Jasper County. The entire county. Jeff Davis County. The entire county. Jefferson County. The entire county. Jenkins County. The entire county. Johnson County. The entire county. Jones County. The entire county. Lamar County. The entire county. Lanier County. The entire county. Laurens County. The entire county. Lee County. The entire county. Liberty County. The entire county. Lincoln County. The entire county. Long County. The entire county.

Lowndes County. The entire county. Macon County. The entire county. Madison County. The entire county.

Marion County. The entire county. McDuffie County. The entire county. McIntosh County. The entire county. Meriwether County. The entire county. Miller County. The entire county. Mitchell County. The entire county. Monroe County. The entire county. Montgomery County. The entire county.

Morgan County. The entire county. Muscogee County. The entire county. Newton County. The entire county. Oconee County. The entire county. Oglethorpe County. The entire county. Paulding County. The entire county. Peach County. The entire county. Pierce County. The entire county. Pike County. The entire county. Polk County. The entire county. Pulaski County. The entire county. Putnam County. The entire county. Quitmon County. The entire county. Randolph County. The entire county. Richmond County. The entire county. Rockdale County. The entire county. Schley County. The entire county. Screven County. The entire county. Seminole County. The entire county. Spalding County. The entire county. Stewart County. The entire county. Sumter County. The entire county. Talbot County. The entire county. Taliaferro County. The entire county. Tattnall County. The entire county. Taylor County. The entire county. Telfair County. The entire county.

Terrell County. The entire county.
Thomas County. The entire county.
Tift County. The entire county.
Toombs County. The entire county.
Treutlen County. The entire county.
Troup County. The entire county.
Truner County. The entire county.
Twiggs County. The entire county.
Upson County. The entire county.
Walton County. The entire county.
Waren County. The entire county.
Warnen County. The entire county.
Washington County. The entire county.
Wayne County. The entire county.
Webster County. The entire county.
Wheeler County. The entire county.
Whitfield County. That portion of the county within Georgia Militia Districts 627,

872, 1233, 1298, 1305, and 1433.

Wilcox County. The entire county.

Wilkes County. The entire county.

Wilkinson County. The entire county.

Worth County. The entire county.

(2) Suppressive areas. None.

Louisiana

(1) Generally infested areas. The entire state.

(2) Suppressive areas. None.

Mississippi

(1) Generally infested areas.

Adams County. The entire county.

Alcorn County. The entire county.

Amite County. The entire county.

Attala County. The entire county.

Benton County. That portion of the county south of the north line of T. 4 S.

Bolivar County. T. 20 N., R. 8, 7, and 8 W.;

T. 21 N., R. 5, 6, and 7 W., and S. ½ T. 22 N.,

R. 6 W.

Carroll County. The entire county. Calhoun County. The entire county. Chickasaw County. The entire county. Choctaw County. The entire county. Claiborne County. The entire county. Clarke County. The entire county. Clay County. The entire county. Copiah County. The entire county. Covington County. The entire county. Forrest County. The entire county. Franklin County. The entire county. George County. The entire county. Greene County. The entire county. Grenada County. The entire county. Hancock County. The entire county. Harrison County. The entire county. Hinds County. The entire county. Holmes County. The entire county. Humphreys County. The entire county. Issaquena County. The entire county. Itawamba County. The entire county. Jackson County. The entire county. Jasper County. The entire county. Jefferson County. The entire county. Jefferson Davis County. The entire county. Jones County. The entire county. Kemper County. The entire county. Lafayette County. That portion of the county south of the north line of T. 10 S.; T. 9

W.

Lamar County. The entire county.

Lauderdale County. The entire county.

Lawrence County. The entire county.

S., R. 1, 2, 3, and 4, W.; T. 8 S., R. 1, 2, and 3, W.; T. 7 S., R. 1 W., and S.E. ¼, T. 6 S., R. 3

Leake County. The entire county.
Lee County. The entire county.
Leflore County. That portion of the county south of the north line of T. 19 N.; S. ½ of T. 20 N., R. 1 E.; and that portion of T. 20 and 21 N., R. 2 E. within the county.

Lincoln County. The entire county.
Lowndes County. The entire county.
Madison County. The entire county.
Marion County. The entire county.
Marshall County. That portion of T. 7 S., R.

Marshall County. That portion of 1.7 S., R.

1 W. and T. 6 S., R. 1 W. within the county.

Monroe County. The entire county.

Montgomery County. The entire county.

Neshoba County. The entire county.

Newton County. The entire county.

Noxubee County. The entire county.

Oktibeha County. The entire county.

Panola County. That portion of T. 10 S., R. 5

W. within the county and T. 10 S., R. 6 and 7

W.

Pearl River County. The entire county.
Perry County. The entire county.
Pike County. The entire county.
Pontotoc County. The entire county.
Prentiss County. The entire county.
Rankin County. The entire county.
Scott County. The entire county.
Sharkey County. The entire county.
Simpson County. The entire county.
Smith County. The entire county.
Stone County. The entire county.
Sunflower County. That portion of the county south of the north line of T. 19 and 20 N., R. 5 W.

Tallahatchie County. That portion of the county south of the north line of T. 24 N., and east of the west line of R. 2 E., and T. 25 N., R. 3 E.

Tippah County. The entire county.
Tishomingo County. The entire county.
Union County. The entire county.
Walthall County. The entire county.
Warren County. The entire county.
Washington County. The entire county.
Wayne County. The entire county.
Webster County. The entire county.
Wilkinson County. The entire county.
Winston County. The entire county.
Yalobusha County. The entire county.
Yazoo County. The entire county.
(2) Suppressive areas. None.

North Carolina

(1) Generally infested areas. Anson County. That portion of the county bounded by a line beginning at the intersection of State Secondary Road 1756 and the Pee Dee River; then southwesterly along this road to its intersection with State Secondary Road 1744; then southerly along this road to its intersection with State Secondary Road 1730; then west along this road to its intersection with State Secondary Road 1801; then southeasterly along this road to its intersection with State Highway 145; then northeasterly along this highway to its intersection with U.S. Highway 74; then east along this highway to its intersection with State Secondary Road 1748; then north along this road to its intersection with the Pee Dee River; then northwesterly along this river to the point of beginning.

Beaufort County. The entire county.

Bladen County. The entire county.

Brunswick County. The entire county.

Carteret County. The entire county.
Columbus County. The entire county.
Craven County. The entire county.

Duplin County. That area bounded by a line beginning at the intersection of State Secondary Road 1130 and the Duplin-Sampson County line; then north along this county line to its intersection with State Secondary Road 1108; then northeasterly along this road to its junction with State Secondary Road 1112; then easterly along this road to its junction with State Secondary Road 1105; then north along this road to its junction with State Secondary Road 1113; then easterly along this road to its junction with State Secondary Road 1106; then south and east along this road to its junction with U.S. Highway 117 and State Secondary Road 1107; then northeast along State Secondary Road 1107 to its intersection with State Secondary Road 1900; then southeast along this road to its junction with State Secondary Road 1003; then east along this road to its intersection with State Highway 11; then northerly along this highway to its junction with State Highway 111; then northerly along this highway to its intersection with State Secondary Road 1555; then northeast along this road to its intersection with State Secondary Road 1553; then southeasterly along this road to its intersection with State Secondary Road 1551; then east along this road to its intersection with State Secondary Road 1549; then southerly along this road to its intersection with State Highway 11; then east along this highway to its intersection with the Duplin-Lenoir County line; then southerly along this county line to its intersection with the Duplin-Jones County line; then southeast along this county line to its intersection with the Duplin-Onslow County line; then southerly along this county line to its intersection with the Duplin-Pender County line; then west along this county line to its intersection with the Duplin-Sampson County line; then westerly along this county line to the point of beginning.

Hyde County. That portion of the county bounded by a line beginning at a point where Scranton Creek intersects U.S. Highway 264; then north along this highway to its junction with State Secondary Road 1302; then north along this road to its junction with State Secondary Road 1303; then northeast along this road to its intersection with the northwest fork of New Lake Fork Creek; then east along this creek to its junction with Alligator River; then east along this river to its intersection with State Highway 94; then south along this highway to its junction with State Secondary Road 1311; then easterly along this road to its junction with U.S. Highway 264; then southwest along this highway to its junction with State Secondary Road 1164; then southeast along this road to its junction with the Pamlico Sound; then westerly along the mainland Hyde County-Pamlico Sound shoreline to its junction with the Pungo River; then northerly along this river to its junction with Scranton Creek; then southeasterly along this creek to the point of beginning.

eginning.

Jones County. The entire county.

Lenoir County. That portion of the county bounded by a line beginning at a point where State Secondary Road 1165 intersects the Duplin-Lenoir County line; then east along this road to its junction with State Highway 11; then northerly along this highway to its intersection with U.S. Highway 70 (bypass); then easterly along this highway to its intersection with the Jones-Lenoir County line; then southerly along this county line to the Duplin-Lenoir County line; then northerly along this county line to the point of beginning.

New Hanover County. The entire county.
Onslow County. The entire county.
Pamlico County. The entire county.
Pender County. The entire county.

Richmond County. Beginning at the junction of the Pee Dee River and U.S. Highway 74, then east to the junction of State Road 1140 and U.S. Highway 74; then northeast along State Road 1140 to its junction with State Road 1141; then northwest along this road to its junction with State Road 1144; then northeast along this road to its junction with State Road 1146; then northwest along this road to its junction with State Road 1148; then west along this road to Mountain Creek; then southwest along this creek to the Pee Dee River; then south along this river to the point of beginning.

Robeson County. That portion of the county bounded by a line beginning at the junction of Robeson County and Scotland County at the North Carolina-South Carolina State line; then northeast along the Scotland-Robeson County line to its junction with the Seaboard Coastline Railroad at Maxton, North Carolina; then southeast along this railroad to its intersection of the North-South Seaboard Coastline Railroad at Pembroke, North Carolina; then northeast along this railroad to its junction with North Carolina Highway 211; then southeast along this highway to its junction with State Secondary Road 1529; then northeast along this road to its intersection of U.S. Highway 301; then northeast along this highway to its junction with State Secondary Road 1935; then east by southeast along this road to its intersection of State Secondary Road 1935 and State Secondary Road 1004; then northeast along State Secondary Road 1004 to its junction with the Bladen-Robeson County line; then east and southeast along this county line to the junction of the Bladen-Columbus-Robeson County lines; then south and southwest along the Columbus-Robeson County line to its junction with the North Carolina-South Carolina State line: then northwest along this state line to the point of beginning.

Sampson County. That portion of the county bounded by a line beginning at a point where State Secondary Road 1208 intersects the Sampson-Bladen County line; then northeast along this road to its junction with State Secondary Road 1210; then north along this road to its junction with State Highway 411; then north along this highway to its intersection with State Secondary Road 1214; then northeast along this road to its junction with State Secondary Road 1219; then easterly along this road to its junction with U.S. Highway 701; then south along this highway to its junction with State Secondary Road 1145; then easterly along this road to its junction with State Secondary Road 1147;

then north and east along this road to its junction with State Secondary Road 1938; then easterly along this road to its junction with State Secondary Road 1004; then northeasterly along this road to its intersection with State Secondary Road 1926; then east along this road to its intersection with the Sampson-Duplin County line; then southerly and easterly along this county line to the Sampson-Pender County line; then southwesterly along this county line to the Sampson-Bladen County line; then northwesterly along this county line to the point of beginning.

Scotland County. Beginning at the junction of the North Carolina-South Carolina State line and State Road 1619; then northeast along this road to its junction with U.S. Highway 501; then northwest along this highway to its junction with U.S. 401 Business: then northeast along this highway to the Hoke-Scotland County line and junction with the Lumber River; then southeast along this river to its junction with the Robeson-Scotland County line; then southwest along this county line to the North Carolina-South Carolina State line; then northwest along this state line to the point of beginning.

Union County. Beginning at a point where U.S. Highway 74 intersects the Union-Anson County line; then south along this county line to its junction with the North Carolina-South Carolina State line; then west along this state line to its junction with the Lancaster County line; then north and northwest along this county line to its intersection with the Mecklenburg-Union County line; then northeast along this county line to its intersection with U.S. Highway 74; then southeast and east along this highway to the point of beginning.

(2) Suppressive areas. None.

Oklahoma

(1) Generally infested areas.

Bryan County. That portion of the county south of the north line of T. 5 S., R. 7, 8, and 9 F.

Marshall County. That portion of the county south of the north line of T. 6 S., R. 6, and 7 E.

McCurtain County. That portion of the county south of the north line of T. 7 S., R. 21, 22, 23, 24, 25, 26, and 27 E.

(2) Suppressive areas. None.

South Carolina

(1) Generally infested areas.

Aiken County. The entire county.

Allendale County. The entire county.

Bamberg County. The entire county.

Barnwell County. The entire county.

Beaufort County. The entire county.

Berkeley County. The entire county.

Calhoun County. The entire county.

Charleston County. The entire county.

Chesterfield County. The entire county.

Clarendon County. The entire county.

Chester County. That portion of the county bounded by a line beginning at a point where State Primary Highway 72 intersects with the Chester-Union County line; then northeast along this highway to its junction with State Primary Highway 9; then easterly along this highway to its intersection with the Chester-

Lancaster County line; then south along this county line to its junction with the Chester-Fairfield County line; then west along this county line to the Union County line; then north along this county line to the point of beginning.

Colleton County. The entire county. Darlington County. The entire county. Dillon County. The entire county. Dorchester County. The entire county. Edgefield County. The entire county. Fairfield County. The entire county. Florence County. The entire county. Georgetown County. The entire county. Hampton County. The entire county. Horry County. The entire county. Jasper County. The entire county. Kershaw County. The entire county. Lancaster County. The entire county. Lee County. The entire county. Lexington County. The entire county. Marion County. The entire county. Marlboro County. The entire county

McCormick County. That portion of the county bounded by a line beginning at a point where U.S. Highway 378 junctions with the Clark Hill Reservoir; then northeast along this highway to its intersection with the McCormick-Edgefield County line; then southerly along this county line to its junction with the Savannah River; then northwesterly along this river and Clark Hill Reservoir to the point of beginning.

Newberry County. That portion of the county bounded by a line beginning at a point where U.S. Highway 76 intersects with the Newberry-Laurens County line; then northeasterly, easterly, southerly, and westerly along the Newberry County line to its intersection with State Primary Highway 121; then north along this highway to its junction with State Primary Highway 34; then northeast along this highway to its junction with U.S. Highway 76; then northwest along this highway to the point of beginning.

Richland County. The entire county.
Saluda County. The entire county.
Sumter County. The entire county.
Union County. That portion of the county bounded by a line beginning at a point where State Primary Highway 72 intersects with the Union-Newberry County line; then northeast along this highway to its junction with the Broad River; then south along this river to its junction with the Newberry County line; then west and north along this county line to the point of beginning

Orangeburg County. The entire county.

point of beginning.

Williamsburg County. The entire county.

[2] Suppressive areas. None.

Texas

(1) Generally infested areas.
Anderson County. The entire county.
Angelina County. The entire county.
Aransas County. The entire county.
Atascosa County. The entire county.
Austin County. The entire county.
Bandera County. The entire county.
Bestrop County. The entire county.
Beel County. The entire county.
Beell County. The entire county.
Bexar County. The entire county.
Blanco County. The entire county.
Bosque County. The entire county.
Bowie County. The entire county.

Brazoria County. The entire county. Brazos County. The entire county. Burleson County. The entire county. Burnet County. The entire county. Caldwell County. The entire county. Calhoun County. The entire county. Camp County. The entire county. Cass County. The entire county. Chambers County. The entire county. Cherokee County. The entire county. Collin County. The entire county. Colorado County. The entire county. Comal County. The entire county. Comanche County. The entire county.
Cooke County. The entire county. Coryell County. The entire county. Dallas County. The entire county. Denton County. The entire county. De Witt County. The entire county. Duval County. That portion of the county within a 3 mile radius of the intersection of

state highway 44 and state highway 359. Eastland County. The entire county. Ellis County. The entire county. Erath County. The entire county. Falls County. The entire county. Fannin County. The entire county. Fayette County. The entire county. Fort Bend County. The entire county. Franklin County. The entire county. Freestone County. The entire county. Frio County. The entire county. Galveston County. The entire county.
Gillespie County. The entire county.
Goliad County. The entire county. Gonzales County. The entire county. Grayson County. The entire county. Gregg County. The entire county. Grimes County. The entire county. Guadalupe County. The entire county. Hardin County. The entire county. Harris County. The entire county. Harrison County. The entire county. Hays County. The entire county. Henderson County. The entire county. Hill County. The entire county. Hood County. The entire county. Houston County. The entire county. Hunt County. The entire county. Jackson County. The entire county. Jasper County. The entire county. Jefferson County. The entire county. Jim Wells County. The entire county. Johnson County. The entire county. Karnes County. The entire county. Kaufman County. The entire county. Kendall County. The entire county. Kerr County. The entire county

Kimble County. The portion of the county bounded by a line beginning at a point where Texas Ranch Road 479 intersects the Kerr-Kimble County line, then northerly along this road to its junction with U.S. Highway 290, then easterly along this highway to its intersection with the Gillespie-Kimble County line, then southerly along this county line to its intersection with the Kerr-Gillespie County line; then westerly along this county line to the point of beginning.

Kleberg County. The entire county.
Lampasas County. The entire county.
Lavaca County. The entire county.
Lee County. The entire county.
Leon County. The entire county.
Liberty County. The entire county.

Limestone County. The entire county.

Live Oak County. The entire county.

Llano County. The portion of the county bounded by a line beginning where Texas Ranch 965 intersects the Llano-Gillespie County line; then northeasterly along this road to its intersection with Texas Highway 16; then northerly along this highway to its intersection with Texas Highway 71; then southeasterly along this highway to its intersection with Texas Ranch Road 2233; then northeasterly along this road to its intersection with the Burnet-Llano County line; then southeasterly along this county line to its junction with the Blanco-Burnet-Llano County line; then westerly along this county line to the point of beginning.

Madison County. The entire county.

Marion County. The entire county. Matagorda County. The entire county. McLennan County. The entire county. Medina County. The entire county.

Milan County. The entire county.

Montgomery County. The entire county. Morris County. The entire county. Nacogdoches County. The entire county. Navarro County. The entire county. Newton County. The entire county. Nueces County. The entire county. Orange County. The entire county. Panola County. The entire county.
Parker County. The entire county. Polk County. The entire county. Rains County. The entire county. Refugio County. The entire county. Robertson County. The entire county. Rockwall County. The entire county. Rusk County. The entire county. Sabine County. The entire county. San Augustine County. The entire county. San Jacinto County. The entire county. San Patricio County. The entire county. Shelby County. The entire county. Smith County. The entire county. Somervell County. The entire county. Tarrant County. The entire county. Taylor County. The entire county. Titus County. The entire county. Tom Green County. The entire county. Travis County. The entire county. Trinity County. The entire county. Tyler County. The entire county. Upshur County. The entire county.

Uvalde County. The portion of the county bounded by a line beginning at a point where U.S. Highway 90 intersects the Medina-Uvalde County line; then westerly along this highway to its intersection with Texas Ranch Road 2730; then northeasterly along this road to its intersection with Texas Highway 127; then northwesterly along this highway to its intersection with U.S. Highway 83; then northerly along this highway to its junction with the Real-Uvalde County line; then easterly along this county line to its junction with the Bandera-Uvalde County line; then easterly along this county line to its junction with the Medina-Uvalde County line; then southerly along this county line to the point of beginning.

Van Zandt County. The entire county.
Victoria County. The entire county.
Walker County. The entire county.
Waller County. The entire county.
Washington County. The entire county.
Wharton County. The entire county.

Wichita County. The entire county.
Williamson County. The entire county.
Wilson County. The entire county.
Wise County. The entire county.
Wood County. The entire county.
Young County. Those portions of the county within a 3 mile radius from the intersection of Farm to Market Road 1287 and state highway 16, and within a 3 mile radius from the intersection of Farm to Market Road 210 and state highway Spur 132.

(2) Suppressive areas. None. Done at Washington, DC, this 26th day of August, 1987.

D. Husnik,

Acting Deputy Administrator, Plant Protection and Quarantine, Animal and Plant Health Inspection Service. [FR Doc. 87–20078 Filed 8–31–87; 8:45 am] BILLING CODE 3410–34-M

DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Part 39

[Docket No. 87-CE-26-AD; Amdt. 39-5709]

Airworthiness Directives; Beech Aircraft Corporation Model 58 Airplanes

AGENCY: Federal Aviation Administration (FAA), DOT. ACTION: Final rule.

SUMMARY: This amendment adopts a new Airworthiness Directive (AD), applicable to certain Beech Aircraft Corporation Model 58 airplanes which are approved for flight in icing conditions. The FAA has determined that the heating circuitry for the stall warning vane on these airplanes will not provide sufficient heat for flight in icing conditions. This AD requires installation of a placard, "Flight into Known Icing Conditions Prohibited," and a copy of the AD placed in the Limitations Section of Pilot's Operating Handbook and FAA Approved Airplane Flight Manual, unless it is determined by an inspection that a relay is installed in the stall warning vane heater circuit. The placard and limitation may be removed after the relay is installed. The actions of this AD are necessary to preclude loss of stall warning capability in icing conditions.

DATES: Effective Date: September 1, 1987

Compliance: As prescribed in the body of the AD.

ADDRESSES: Beechcraft Mandatory Service Bulletin (MSB) No. 2180 dated June 1987, applicable to this AD may be obtained from Beech Aircraft Corporation, Commercial Service, Department 52, P.O. Box 85, Wichita, Kansas 67201–0085. This information may be examined at the Rules Docket, Office of the Regional Counsel, Room 1558, 601 East 12th Street, Kansas City, Missouri 64106.

FOR FURTHER INFORMATION CONTACT: Dale A. Vassalli, Aerospace Engineer, ACE-130W, Wichita Aircraft Certification Office, 1801 Airport Road, Room 100, Mid-Continent Airport, Wichita, Kansas 67209; Telephone (316) 946-4419.

SUPPLEMENTARY INFORMATION: Beech has made a production change on certain Model 58 airplanes, which included relocation and use of a different relay in the stall warning vane heater circuit. This production change, on airplanes which are approved for flight in icing conditions, provides sufficient heat for the stall warning vane for correct operation in icing conditions. It has been subsequently determined that some of these airplanes did not have the relay installed during production. Without the installation of the relay, sufficient heat will not be applied to the stall warning vane, thereby jeopardizing its stall warning capability and creating a potentially unsafe condition.

As a result, Beech issued MSB No. 2180, dated June 1987, which defines a procedure for determining if the relay is installed, and instructions for installation of the relay if it is missing. Beech has recommended compliance with the Service Bulletin within the next 100 hours time-in-service. The FAA is in agreement with the inspection and modification compliance time; however, since a heated stall warning system is required for flight in icing conditions, the affected airplanes will be prohibited from flight into icing conditions until the actions specified in MSB No. 2180 have been accomplished.

Since the FAA has determined that the unsafe condition described herein is likely to exist in other airplanes of the same type design, an AD is being issued, applicable to certain Beech Model 58 airplanes, which prohibits operation in icing conditions until the airplane is inspected and modified as required. Because an emergency condition exists that requires the immediate adoption of this regulation, it is found that notice and public procedure hereon are impractical and contrary to the public interest, and good cause exists for making this amendment effective in less than 30 days.

The FAA has determined that this regulation is an emergency regulation that is not major under section 8 of Executive Order 12291. It is impracticable for the agency to follow

the procedures of Order 12291 with respect to this rule since the rule must be issued immediately to correct an unsafe conditions in aircraft. It has been further determined that this document involves an emergency regulation under DOT Regulatory Policies and Procedures (44 FR 11034; February 26, 1979). If this action is subsequently determined to involve a significant regulation, a final regulatory evaluation or analysis, as appropriate, will be prepared and placed in the regulatory docket (otherwise, an evaluation is not required). A copy of it, when filed, may be obtained by contacting the Rules Docket under the caption "Addresses" at the location identified.

List of Subjects in 14 CFR Part 39

Air transportation, Aviation safety, Aircraft, Safety.

Adoption of the Amendment

Accordingly, pursuant to the authority delegated to me by the Administrator, the Federal Aviation Administration amends § 39.13 of Part 39 of the FAR as follows:

PART 39-[AMENDED]

1. The authority citation for Part 39 continues to read as follows:

Authority: 49 U.S.C. 1354(a), 1421 and 1423; 49 U.S.C. 106(g) (Revised, Pub. L. 97–449, January 12, 1983); and 14 CFR 11.89.

§ 39.13 [Amended]

2. By adding the following new AD:

Beech Aircraft Corporation: Applies to Model 58 (S/N's TH-1389, TH-1396, TH-1397, TH-1403, TH-1407 through TH-1410, TH-1412, TH-1414, TH-1416, TH-1420 through TH-1423, TH-1425, TH-1427, TH-1430, TH-1434, TH-1438, TH-1440, TH-1442, TH-1434, TH-1445, TH-1445, TH-1451, TH-1452, TH-1456, TH-1457, TH-1459, TH-1462, TH-1463, TH-1466, TH-1468, TH-1477, TH-1478, TH-1472 through TH-1474, TH-1477, TH-1478, TH-1481, TH-1483, TH-1485, TH-1490, TH-1492, TH-1495 through TH-1497, TH-1493, TH-1495 through TH-1497, TH-1499, TH-1500, TH-1502 and TH-1506) airplanes certificated in any category.

Compliance: Required as indicated after the effective date of this AD, unless

previously accomplished.

To preclude the loss of stall warning capability in icing conditions, accomplish the following:

- (a) Within the next 25 hours time-in-service (unless the inspection and relay installation requirements of paragraph (c) have been accomplished):
- (1) Fabricate and install on the instrument panel in clear view of the pilot the following placard using letters of a minimum of 0.10 inch in height:

"FLIGHT INTO KNOWN ICING CONDITIONS IS PROHIBITED".

(2) Place a copy of this AD in the Limitation Section of Pilot's Operating Handbook (POH) and FAA Approved Airplane Flight Manual (AFM).

(3) Cover the airplane operating placard statement, "This airplane approved for flight in icing conditions" with opaque tape.

(4) Operate the airplane in accordance with

this limitation.

(b) The requirements of paragraph (a) of this AD may be accomplished by the owner/ operator on any airplanes which are not used under FAR Part 121 or 135. The person accomplishing these actions must make the appropriate airplane maintenance record entry per FAR 43.9 and 91.173.

(c) Within the next 100 hours time-inservice after the effective date of this AD, inspect and, if necessary, modify the stall warning heater circuit in accordance with the instructions in Beechcraft Mandatory Service Bulletin No. 2180, dated June 1987.

(d) The requirements and limitations of paragraph (a) of this AD do not apply to airplanes that have been inspected and modified as required per paragraph (c) of this AD.

(e) Airplanes may be flown in accordance with FAR 21.197 to a location where this AD

may be accomplished.

(f) An equivalent means of compliance with this AD may be used if approved by the Manager, Wichita Aircraft Certification Office, 1801 Airport Road, Room 100, Mid-Continent Airport, Wichita, Kansas 67209; Telephone (316) 946–4400.

All persons affected by this directive may obtain copies of the document(s) referred to herein upon request to Beech Aircraft Corporation, Commercial Service, Department 52, P.O. Box 85, Wichita, Kansas 67201–0085; or may examine the document(s) referred to herein at FAA, Office of the Regional Counsel, Room 1558, 601 East 12th Street, Kansas City, Missouri 64106.

This amendment becomes effective on September 1, 1987.

Issued in Kansas City, Missouri, on August 7, 1987.

Berry D. Clements,

Acting Director, Central Region.
[FR Doc. 87–19997 Filed 8–31–87; 8:45 am]
BILLING CODE 4910–13–M

14 CFR Part 39

[Docket No. 87-ASW-24; Amdt. 39-5678]

Airworthiness Directives; McDonnell Douglas Helicopter Company, Model 369D, E, F, and FF Helicopters

AGENCY: Federal Aviation Administration (FAA), DOT. ACTION: Final rule.

SUMMARY: This amendment adopts a new airworthiness directive (AD) which requires inspection of the main rotor transmission tail rotor output drive pinion shaft and removal from service of all unairworthy shafts on McDonnell Douglas Helicopter Company (MDHC) Model 369D, E, F, and FF helicopters. The AD is prompted by reports that main rotor transmission tail rotor output drive shafts have failed in flight, which could result in loss of power to the tail rotor and loss of control of the helicopter.

DATES: September 1, 1987.

Effective Date: The incorporation by reference of certain publications listed in the regulations is approved by the Director of the Federal Register as of September 1, 1987.

Compliance: As indicated in the body of this AD.

ADDRESSES: The applicable service information notice may be obtained from McDonnell Douglas Helicopter Company, 500 E. McDowell Road, Mesa, Arizona 85205

A copy of each document supporting the AD is contained in the Rules Docket, Office of the Regional Counsel, Federal Aviation Administration, Southwest Region, Room 158, Building 3B, 4400 Blue Mound Road, Fort Worth, Texas.

FOR FURTHER INFORMATION CONTACT:
Mr. William R. Twa, Jr., Aerospace
Engineer, Propulsion Section, ANM—
174W, Western Aircraft Certification
Office, Northwest Mountain Region,
Federal Aviation Administration, P.O.
Box 92007, Worldway Postal Center, Los
Angeles, California 90009–2007;
telephone (213) 297–1128.

SUPPLEMENTARY INFORMATION: There have been six reported cases of forward bearing journal failures of the main rotor transmission tail rotor output drive shaft (P/N's 369D25125-BSC and -11) due to fatigue in an area between the forward bearing journal and the toothed gear section. Contributory causes of the fatigue have been traced to the need for a relief radius, omission of shotpeening of the area, improper surface finish, and no plating on the inside diameter of the shaft. All failures have occurred during flight and were detected by a loud noise and vibrations. The currently acceptable nondestructive test method for verifying the integrity of the shaft is a magnetic particle and visual inspection of the affected area. MDHC has issued Mandatory Service Information Notice (SIN) DN-147/EN-35/FN-24, dated April 23, 1987. Since this condition is likely to exist or develop in other helicopters of the same type design, an airworthiness directive is being issued which requires a magnetic particle and visual inspection of the main rotor

transmission tail rotor output drive pinion shaft (P/N's 369D25125–BSC and -11), and removal of unairworthy pinion shafts from service on MDHC Model 369D, E, F, and FF helicopters.

Since a situation exists that requires immediate adoption of this regulation, it is found that notice and public procedure hereon are impracticable and good cause exists for making this amendment effective in less than 30 days.

The FAA has determined that this regulation is an emergency regulation that is not considered to be major under Executive Order 12291. It is impracticable for the agency to follow the procedures of Order 12291 with respect to this rule since the rule must be issued immediately to correct an unsafe condition in aircraft. It has been further determined that this action involves an emergency regulation under DOT Regulatory Polices and Procedures (44 FR 11034; February 26, 1979). If this action is subsequently determined to involve a significant/major regulation, a final regulatory evaluation or analysis, as appropriate, will be prepared and placed in the regulatory docket (otherwise, an evaluation or analysis is not required). A copy of it, when filed, may be obtained by contacting the person identified under the caption "FOR **FURTHER INFORMATION CONTACT."**

List of Subjects in 14 CFR Part 39

Air transportation, Aircraft, Aviation safety, Safety and Incorporation by reference.

Adoption of the Amendment

Accordingly, pursuant to the authority delegated to me by the Administrator, the Federal Aviation Administration amends § 39.13 of Part 39 of the Federal Aviation Regulations as follows:

PART 39—AIRWORTHINESS DIRECTIVES

1. The authority citation for Part 39 continues to read as follows:

Authority: 49 U.S.C. 1354(a), 1421 and 1423; 49 U.S.C. 106(g) (Revised, Pub. L. 97–449, January 12, 1983); and 14 CFR 11.89.

§ 39.13 [Amended]

2. By adding the following new AD:

McDonnell Douglas Helicopter Company (MDHC) (Hughes Helicopters, Inc.):
Applies to all Model 369D, E, F, and FF helicopters, certificated in any category, equipped with MDHC main transmission (P/N 369D25100-BSC or -501) which has tail rotor output drive pinion shaft (P/N 369D25125-BSC or -11) installed.

Compliance is required as indicated, unless previously accomplished.

To prevent possible loss of tail rotor control, accomplish the following: (a) Within the next 25 hours' time in service after the effective date of this AD, perform a one-time magnetic particle and visual inspection on the main transmission tail rotor output drive pinion shaft (P/N 369D25125-BSC or -11) on helicopters with the following serialized main rotor transmission in accordance with procedures detailed in paragraphs (c) through (f) of this AD.

Transmission Serial Number

1989 1992 and 1993 1998 through 2000 2002 through 2082 2084 and 2085

If the tail rotor output drive pinion shaft serial number is contained in the list below, remove the shaft from service prior to further flight

Tail Roter Output Drive Pinion Shaft Serial

Numbers 1474 through 1502 1504 through 1547 1549 through 1561 1563 and 1564 1666

(b) Within the next 100 hours' time in service after the effective date of this AD, perform a one-time magnetic particle and visual inspection on the main transmission tail rotor output drive pinion shaft (P/N 369D25125-BSC or -11) on all affected helicopters with transmission serial numbers other than those listed above in accordance with procedures detailed in paragraphs (c) through (f) of this AD.

(c) Verify that the tail rotor output drive pinion shaft has an undercut style fillet radius as shown in Figure 1 of MDHC Mandatory SIN DN-147/EN-35/FN-24, dated April 23, 1987, or FAA-approved equivalent. Remove parts which do not have the undercut from service prior to further flight.

Note.—Tail rotor output drive pinion shafts removed from service in accordance with paragraph (c) may be returned to MDHC for rework if a magnetic particle inspection of the part does not show any crack indications.

(d) Perform a magnetic particle inspection of the tail rotor output drive pinion shaft. Remove parts which show indications of cracking found during magnetic particle inspection from service prior to further flight.

(e) Inspect the tail rotor output drive pinion shafts for scratches, tooling marks, corrosion, or other minor surface defects in the fillet radius or elsewhere on the shaft in accordance with paragraphs c.(4), c.(5), and c.(6) of MDHC Mandatory SIN DN-147/EN-35/FN-24, dated April 23, 1987, Prior to further flight, rework discrepant shafts that show no indication of cracks in accordance with procedures stated in paragraph d. of MDHC Mandatory SIN DN-147/EN-35/FN-24, dated April 23, 1987.

(f) Special flight permits may be issued in accordance with FAR §§ 21.197 and 21.199 to ferry aircraft to a maintenance base in order to comply with the requirements of this AD.

The procedure shall be done in accordance with MDHC SIN DN-147/EN-35/FN-24, dated April 23, 1987. This incorporation by reference was approved by the Director of the Federal Register in accordance with 5 U.S.C. 552(a)(1). Copies may be obtained from MDHC, 500 E. McDowell Road, Mesa, Arizona. Copies may be inspected at the Office of the Regional Counsel, FAA, Southwest Region, 4400 Blue Mound Road, Fort Worth, Texas, or at the Office of the Federal Register, 1100 L. Street, NW., Room 8401, Washington, DC.

This amendment becomes effective September 1, 1987.

Issued in Fort Worth, Texas, on July 9, 1987.

Don P. Watson,

Acting Director, Southwest Region.
[FR Doc. 87–19996 Filed 8–31–87; 8:45 am]
BILLING CODE 4910–13-W

14 CFR Part 71

[Airspace Docket No. 87-ANM-9]

Establish Transition Area, Gooding, ID

AGENCY: Federal Aviation Administration (FAA), DOT.

ACTION: Final rule.

summary: This action etablishes a 700 foot transition area at Gooding, Idaho, to provide controlled airspace for a nondirectional beacon (NDB) instrument approach procedure for Runway 25 at Gooding, Idaho, Municipal Airport.

EFFECTIVE DATE: 0901 UTC, November 19, 1987.

FOR FURTHER INFORMATION CONTACT: Robert L. Brown, ANM-535, Federal Aviation Administration, Docket No. 87-ANM-9, 17900 Pacific Highway South, C-68966, Seattle, Washington 98168, Telephone: (206) 431-2535.

SUPPLEMENTARY INFORMATION:

History

On June 22, 1987, the FAA proposed to amend Part 71 of the Federal Aviation Regulations (14 CFR Part 71) to establish a 700 foot transition area at Gooding, Idaho (52 FR 23468).

Interested parties were invited to participate in this rulemaking proceeding by submitting written comments on the proposal at the FAA. No comments objecting to the proposal were received. Except for editorial changes, this amendment is the same as that proposed in the notice. Section 71.181 of Part 71 of the Federal Aviation

Regulations was republished in Handbook.7400.6C dated January 2, 1987.

The Rule

This amendment to Part 71 of the Federal Aviation Regulations establishes a 700 foot transition area to provide controlled airspace for aircraft executing a new standard instrument approach procedure to the Gooding Municipal Airport.

The FAA has determined that this proposed regulation only involves an established body of technical regulations for which frequent and routine amendments are necessary to keep them operationally current. It, therefore-(1) is not a "major rule" under Executive Order 12291; [2] is not a "significant rule" under DOT Regulatory Policies and Procedures (44 FR 11034; February 26, 1979); and (3) does not warrant preparation of a regulatory evaluation as the anticipated impact is so minimal. Since this is a routine matter that will only affect air traffic procedures and air navigation, it is certified that this rule, when promulgated, will not have a significant economic impact on a substantial number of small entities under the criteria of the Regulatory Flexibility Act.

List of Subjects in 14 CFR Part 71

Aviation safety, Transition areas. Adoption of the Amendment

Accordingly, pursuant to the authority delegated to me, Part 71 of the Federal Aviation Regulations (14 CFR Part 71) is amended as follows:

PART 71-[AMENDED]

1. The authority citation for Part 71 continues to read as follows:

Authority: 49 U.S.C. 1348(a), 1354(a), 1510; E.O. 10854; 49 U.S.C. 106(g) (Revised Pub. L. 97–449, January 12, 1983); 14 CFR 11.69.

§ 71.181 [Amended]

2. Section 71.181 is amended as follows:

Gooding, Idaho, Transition Area [New]

That airspace extending upward from 700 feet above the surface within a 9.5 mile radius of the Gooding, Idaho, Municipal Airport [lat. 42°54'45" W., long. 114°45'50" W.

Issued in Seattle, Washington, on August 17, 1987.

Temple H. Johnson, Jr.,

Manager, Air Traffic Division, Northwest Mountain Region

[FR Doc. 87-19992 Filed 8-31-87; 8:45 am] BILLING CODE 4910-13-M

14 CFR Part 71

[Airspace Docket No. 87-ANM-13]

Alteration of Transition Area, Glendive, MT

AGENCY: Federal Aviation Administration (FAA), DOT. ACTION: Final rule.

SUMMARY: This action amends the Glendive, Montana, transition area by adding 1,200 foot transition airspace to the existing transition area description. This change has no effect on the existing 700 foot transition area.

EFFECTIVE DATE: 0901 UTC, October 10, 1987.

FOR FURTHER INFORMATION CONTACT: Robert L. Brown, ANM-535, Federal Aviation Administration, Docket No. 87– ANM-13, 17900 Pacific Highway South, C-68966, Seattle, Washington 98168, Telephone: (206) 431–2535.

SUPPLEMENTARY INFORMATION:

History

On June 22, 1987, the FAA proposed to amend Part 71 of the Federal Aviation Regulations (14 CFR Part 71) to amend the Glendive, Montana, 1,200 foot transition area (52 FR 23467).

Interested parties were invited to participate in this rulemaking proceeding by submitting written comments on the proposal to the FAA. No comments objecting to the proposal were received. Except for editorial changes, this amendment is the same as that proposed in the notice. Section 71.181 of Part 71 of the Federal Aviation Regulations was republished in Handbook 7400.6C dated January 2, 1987.

The Rule

This amendment to Part 71 of the Federal Aviation Regulations amends the Glendive, Montana, 1,200 foot transition area. This change permits the routing of arriving aircraft to the NDB from both Miles City and Williston VORTAC's below 14,500 feet AMSL and allows departures to utilize diverse departure procedures directly to both Williston and Miles City.

The FAA has determined that this proposed regulation only involves an established body of technical regulations for which frequent and routine amendments are necessary to keep them operationally current. It, therefore—(1) is not a "major rule" under Executive Order 12291; (2) is not a "significant rule" under DOT Regulatory Policies and Procedures (44 FR 11034; February 26, 1979); and (3) does not warrant preparation of a regulatory

evaluation as the anticipated impact is so minimal. Since this is a routine matter that will only affect air traffic procedures and air navigation, it is certified that this rule, when promulgated, will not have a significant economic impact on a substantial number of small entities under the criteria of the Regulatory Flexibility Act.

List of Subjects in 14 CFR Part 71

Aviation safety, Transition areas.

Adoption of the Amendment

Accordingly, pursuant to the authority delegated to me, Part 71 of the Federal Aviation Regulations (14 CFR Part 71) is amended as follows:

PART 71-[AMENDED]

1. The authority citation for Part 71 continues to read as follows:

Authority: 49 U.S.C. 1348(a), 1354(a), 1510; E. O. 10854; 49 U.S.C. 106(g) (Revised Pub. L. 97–449, January 12, 1983); 14 CFR 11.69.

§ 71.181 [Amended]

2. Section 71.181 is amended as follows:

Glendive, Montana, Transition Area [Amended]

After the words . . . "to 18½ miles northwest of the airport"; add the words, "and that airspace extending upward from 1,200 feet above the surface bounded on the east and southeast by the west edge of V-545 and on the northwest by the east edge of V-465."

Issued in Seattle, Washington, on August 17, 1987.

Temple H. Johnson, Jr.,

Manager, Air Traffic Division, Northwest Mountain Region.

[FR Doc. 87-19993 Filed 8-31-87; 8:45 am] BILLING CODE 4910-13-M

14 CFR Part 71

[Airspace Docket No. 87-ANM-7]

Alteration of Transition Area, Rock Springs, WY

AGENCY: Federal Aviation Administration (FAA), DOT.

ACTION: Final rule.

SUMMARY: This action will alter the Rock Springs, Wyoming, transition area to provide additional controlled airspace east of Rock Springs.

EFFECTIVE DATE: 0901 UTC, November 19, 1987.

FOR FURTHER INFORMATION CONTACT: Robert L. Brown, ANM-535, Federal Aviation Administration, Docket No. 87-ANM-7, 17900 Pacific Highway South,

C-68966, Seattle, Washington 98168, Telephone: (206) 431-2535.

SUPPLEMENTARY INFORMATON:

History

On June 22, 1987, the FAA proposed to amend Part 71 of the Federal Aviation Regulations (14 CFR Part 71) to amend the Rock Springs, Wyoming, transition area (52 FR 23470).

Interested parties were invited to participate in this rulemaking proceeding by submitting written comments on the proposal to the FAA. No comments objecting to the proposal were received. Except for editorial changes, this amendment is the same as that proposed in the notice. Section 71.181 of Part 71 of the Federal Aviation Regulations was republished in Handbook 7400.6C dated January 2, 1987.

The Rule

This amendment to Part 71 of the Federal Aviation Regulations will extend controlled airspace east of Rock Springs, Wyoming, to enable air traffic controllers to radar vector aircraft to the ILS/DME Runway 27 approach to the Sweetwater County Airport. Currently, available airspace is insufficient for this purpose.

The FAA has determined that this proposed regulation only involves an established body of technical regulations for which frequent and routine amendments are necessary to keep them operationally current. It, therefore-(1) is not a "major rule" under Executive Order 12291; (2) is not a "significant rule" under DOT Regulatory Policies and Procedures (44 FR 11034; February 26, 1979); and (3) does not warrant preparation of a regulatory evaluation as the anticipated impact is so minimal. Since this is a routine matter that will only affect air traffic procedures and air navigation, it is certified that this rule, when promulgated, will not have a significant economic impact on a substantial number of small entities under the criteria of the Regulatory Flexibility Act.

List of Subjects in 14 CFR Part 71

Aviation safety, Transition areas.

Adoption of the Amendment

Accordingly, pursuant to the authority delegated to me, Part 71 of the Federal Aviation Regulations (14 CFR Part 71) is amended as follows:

PART 71-[AMENDED]

1. The authority citation for Part 71 continues to read as follows:

Authority: 49 U.S.C. 1348(a), 1354(a), 1510; E.O. 10854; 49 U.S.C. 106(g) (Revised Pub. L. 97-449, January 12, 1983); 14 CFR 11.69.

§ 71.181 [Amended]

2. Section 71.181 is amended as follows:

Rock Springs, Wyoming, Transition Area [Amended]

Change 1,200 foot transition area to read as follows: " * * to 19 miles east of the VORTAC; and that airspace extending upward from 1,200 feet above the surface within a 23-mile radius of the Rock Springs VORTAC, including that airspace bounded by 4.5 miles south of the Rock Springs 099° radial between 23 miles and 42.5 miles, and 4.5 miles east of the Cherokee VORTAC 198° radial between the VORTAC and 56.5 miles, and 4.5 miles northwest of the Rock Springs 051° radial between 23 miles and the Cherokee VORTAC, excluding that airspace included in the Rawlins, Wyoming, transition

Issued in Seattle, Washington, on August 17, 1987.

Temple H. Johnson, Jr.,

Manager, Air Traffic Division, Northwest Mountain Region.

[FR Doc. 87-19991 Filed 8-31-87; 8:45 am] BILLING CODE 4910-13-M

DEPARTMENT OF HEALTH AND **HUMAN SERVICES**

Food and Drug Administration

21 CFR Part 177

[Docket No. 86F-0154]

Indirect Food Additives: Polymers

AGENCY: Food and Drug Administration. ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the food additive regulations to provide for the safe use of ethylene terephthalateisophthalate copolymers that contain at least 97 weight percent of polymer units derived from ethylene terephthalate. This action responds to a petition filed by The Goodyear Tire & Rubber Co. DATES: Effective September 1, 1987;

objections by October 1, 1987. ADDRESS: Written objections to the Dockets Management Branch (HFA-305), Food and Drug Administration, Room 4-62, 5600 Fishers Lane, Rockville,

MD 20857.

FOR FURTHER INFORMATION CONTACT: Hortense S. Macon, Center For Food Safety and Applied Nutrition (HFF-335), Food and Drug Administration, 200 C

Street SW., Washington, DC 20204, 202-

472-5690. SUPPLEMENTARY INFORMATION: In a notice published in the Federal Register

of May 7, 1986 (51 FR 16897), FDA announced that a petition (FAP 5B3884) had been filed by The Goodyear Tire & Rubber Co., 130 Johns Avenue, Akron, OH 44316-0001, proposing that § 177.1630 Polyethylene phthalate polymers (21 CFR 177.1630) be amended to provide for the safe use of ethylene terephthalate-isophthalate copolymers containing a minimum of 97 weight percent of polymer units derived from ethylene terephthalate for use as components of food-contact articles.

FDA has evaluated data in the petition and other relevant materials. The agency concludes that the proposed use of the food additive is safe and that 21 CFR 177.1630(e)(4)(i) should be amended as set forth below.

In accordance with § 171.1(h) (21 CFR 171.1(h)), the petition and the documents that FDA considered and relied upon in reaching its decision to approve the petition are available for inspection at the Center for Food Safety and Applied Nutrition (address above) by appointment with the information contact person listed above. As provided in 21 CFR 171.1(h), the agency will delete from the documents any materials that are not available for public disclosure before making the documents available for inspection.

The agency has carefully considered the potential environmental effects of this action and has concluded that the action will not have a significant impact on the human environment and that an environmental impact statement is not required. The agency's finding of no significant impact and the evidence supporting that finding, contained in an environmental assessment, may be seen in the Dockets Management Branch (address above) betwen 9 a.m. and 4 p.m., Monday through Friday. This action was considered under FDA's final rule implementing the National Environmental Policy Act (21 CFR Part 25)

Any person who will be adversely affected by this regulation may at any time on or before October 1, 1987, file with the Dockets Management Branch (address above) written objections thereto. Each objection shall be separately numbered, and each numbered objection shall specify with particularity the provisions of the regulation to which objection is made and the grounds for the objection. Each numbered objection on which a hearing is requested shall specifically so state. Failure to request a hearing for any particular objection shall constitute a waiver of the right to a hearing on that objection. Each numbered objection for which a hearing is

requested shall include a detailed description and analysis of the specific factual information intended to be presented in support of the objection in the event that a hearing is held. Failure to include such a description and analysis for any particular objection shall constitute a waiver of the right to a hearing on the objection. Three copies of all documents shall be submitted and shall be identified with the docket number found in brackets in the heading of this document. Any objections received in response to the regulation may be seen in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday.

List of Subjects in 21 CFR Part 177

Food additives, Food packaging.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Director and Deputy Director of the Center for Food Safety and Applied Nutrition, Part 177 is amended as follows:

PART 177—INDIRECT FOOD ADDITIVES: POLYMERS

1. The authority citation for 21 CFR Part 177 continues to read as follows:

Authority: Secs. 201(s), 409, 72 Stat. 1784–1788 as amended (21 U.S.C. 321(s), 348); 21 CFR 5.10 and 5.61.

2. In § 177.1630(e)(4)(i) by revising the entry for "Ethylene terephthalate-isophthalate copolymers" to read as follows:

§ 177.1630 Polyethylene phthalate polymers.

(e) * * * (4) * * *

List of Substances and Limitations

(i) Base sheet:

Ethylene terephthalate-isophthalate copolymers: Prepared by the condensation of dimethyl terephthalate or terephthalic acid and dimethyl isophthalate or isophthalic acid with ethylene glycol. The finished copolymers contain either

(a) 77 to 83 weight percent or (b) At least 97 weight percent of polymer units derived from ethylene terephthalate.

Dated: August 24, 1987.

Fred R. Shank,

Acting Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 87-20011 Filed 8-31-87; 8:45 am] BILLING CODE 4160-01-M

21 CFR Parts 510 and 540

Animal Drugs, Feeds, and Related Products; Sterile Procaine Penicillin G Aqueous Suspension (Injectable)

AGENCY: Food and Drug Administration.
ACTION: Final rule.

SUMMARY: The Food and Drug
Administration (FDA) is amending the
animal drug regulations to reflect
approval of a supplemental new animal
drug application (NADA) filed by G.C.
Hanford Manufacturing Co., Inc.,
providing for a reduction of the milk
withholding period after use of sterile
injectable procaine penicillin G aqueous
suspension for treating cattle. In
addition, FDA is amending the firm's
mailing address in the list of sponsors of
approved NADA's to reflect the current
post office box number.

EFFECTIVE DATE: September 1, 1987.

FOR FURTHER INFORMATION CONTACT: Charles E. Haines, Center for Veterinary Medicine (HFV–133), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–443–3410.

SUPPLEMENTARY INFORMATION: G.C. Hanford Manufacturing Co., Inc., P.O. Box 1017, Syracuse, NY 13201, is sponsor of NADA 65-493 which provides for injectable use of sterile procaine penicillin G aqueous suspension to treat cattle, sheep, swine, and horses. The supplement provides for a reduction of the milk withholding period for treating lactating cattle to 48 hours (4 milkings) from 72 hours (6 milkings). The supplement is approved and 21 CFR 540.274b(c) is amended to reflect the approval. The basis for approval is discussed in the freedom of information summary.

In addition, the mailing address for the firm in 21 CFR 510.600(c) is amended to reflect the current post office box number.

In accordance with the freedom of information provisions of Part 20 (21 CFR Part 20) and § 514.11(e)(2)(ii) (21 CFR 514.11(e)(2)(ii), a summary of safety and effectiveness data and information submitted to support approval of this application may be seen in the Dockets Management Branch (HFA-305), Food and Drug Administration, Room 4-62, 5600 Fishers Lane, Rockville, MD 20857, from 9 a.m. to 4 p.m., Monday through Friday.

The Center for Veterinary Medicine has determined pursuant to 21 CFR 25.24(d)(1)(i) that this action is of a type that does not individually or cumulatively have a significant impact on the human environment. Therefore, neither an environmental assessment

nor an environmental impact statement is required.

List of Subjects

21 CFR Part 510

Administrative practice and procedure, Animal drugs, Labeling, Reporting and recordkeeping requirements.

21 CFR Part 540

Animal drugs, Antibiotics.

Therefore, under the Federal Food, Drug, and Cosmetic Act, and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, Parts 510 and 540 are amended as follows:

PART 510-NEW ANIMAL DRUGS

1. The authority citation for 21 CFR Part 510 continues to read as follows:

Authority: Secs. 512, 701(a) (21 U.S.C. 360b, 371(a)); 21 CFR 5.10 and 5.83,

§ 510.600 [Amended]

2. § 510.600 Names, addresses, and drug labeler codes of sponsors of approved applications is amended in paragraph (c)(1) for the entry "G.C. Hanford Manufacturing Co., Inc." and in paragraph (c)(2) for the entry "010515" by revising the post office box number to read "P.O. Box 1017."

PART 540—PENICILLIN ANTIBIOTIC DRUGS FOR ANIMAL USE

3. The authority citation for 21 CFR Part 540 continues to read as follows:

Authority: Sec. 512, 82 Stat. 343-351 (21 U.S.C. 360b); 21 CFR 5.10 and 5.83.

4. § 540.274b is amended in paragraph (c)(3)(ii) by removing the number "010515" and by adding new paragraph (c)(5) to read as follows:

§ 540.274b Procaine penicillin G aqueous suspension.

(c) * * *

(5)(i) Sponsor. See No. 010515 in § 510.600(c) of this chapter.

(ii) See paragraph (c)(3) of this section for specifications, tolerances, and conditions of use of this drug, except that milk taken during treatment and for 48 hours (four milkings) after the latest treatment shall not be used for food.

Dated: August 26, 1987.

Richard A. Carnevale,

Acting Associate Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.

[FR Doc. 87-20012 Filed 8-31-87; 8:45 am] BILLING CODE 4160-01-M

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 52

[A-4-FRL-3254-4; GA-014]

Approval and Promulgation of Implementation Plans; Georgia; PSD Modeling Procedures

AGENCY: Environmental Protection Agency (EPA).

ACTION: Final rule.

SUMMARY: Today, EPA approves a State Implementation Plan revision submitted by the State of Georgia. This revision incorporates regulations promulgated by the Administrator to specify models to be used to comply with the Clean Air Act's requirements for the prevention of significant deterioration (PSD) (sections 165 through 169).

DATES: This action will be effective on November 2, 1987, unless notice is received by October 1, 1987 that someone wishes to submit adverse or critical comments.

ADDRESSES: Copies of the documents relevant to this action are available for public inspection during normal business hours at the following locations:

Public Information Reference Unit, Library Systems Branch, Environmental Protection Agency, 401 M Street, SW., Washington, DC 20460

Environmental Protection Agency, Region IV, Air Programs Branch 345 Courtland Street, NE., Atlanta, GA 30365

Georgia Department of Natural Resources, 205 Butler Street, SE., Floyd Towers East, Atlanta, GA 30334.

FOR FURTHER INFORMATION CONTACT:

Gregg M. Worley, Air Programs Branch, EPA Region IV, at the above address and telephone number (404) 347–2864 or FTS 257–2864.

SUPPLEMENTARY INFORMATION:

Section 165(e)(3)(D) of the Clean Air Act requires the Administrator to promulgate regulations specifying with reasonable particularity models to be used to comply with the Act's PSD requirements. To carry out the requirements, the 1978 "Guideline on Air Quality Models" was incorporated by reference in 40 CFR 51.24 (now renumbered § 51.166) and 40 CFR § 52.21. On September 9, 1986 (51 FR 32176), EPA promulgated amendments to 40 CFR 51.24 (now § 51.166) and 52.21 to substitute by reference the "Guideline on Air Quality Models (Revised)," EPA 450/2-78-027R, in these regulations. This change became effective October 9,

1986. The Clean Air Act, however, gives states nine months (until July 9, 1987 to make the necessary changes in their programs. A review of Georgia's regulations categorized the State as one that does not specifically preclude the use of the revised modeling guideline since 40 CFR 52.21(1) as amended is adopted in the Georgia regulations. The options open to Georgia, therefore, were either to revise their PSD regulation to explicitly include the revised modeling guideline or submit an enforceable letter of commitment in lieu of a regulatory revision. If a letter of commitment is chosen, the letter must mention that the generalized language now means that all PSD permit applicants must use the revised guideline models or models otherwise approved by EPA. Subsequently, Georgia submitted such a letter of commitment on May 11, 1987, with the understanding that it would be processed as a SIP revision.

Final Action

Since Georgia's letter of commitment is consistent with EPA requirements, it is hereby approved. EPA is publishing this action without prior proposal because the Agency views this as a noncontroversial amendment and anticipates no adverse comments. This action will be effective 60 days from the date of this Federal Register unless, within 30 days of its publication, notice is received that adverse or critical comments will be submitted. If such notice is received, this action will be withdrawn before the effective date by publishing two subsequent notices. One notice will withdraw the final action and another will begin a new rulemaking by announcing a proposal of the action and establishing a comment period. If no such comments are received, the public is advised that this action will be effective November 2, 1987.

Under section 307(b)(1) of the Act, petitions for judicial review of this action must be filed in the United States Court of Appeals for appropriate circuit by November 2, 1987. This action may not be challenged later in proceedings to enforce its requirements. (See 307(b)(2).)

Under 5 U.S.C. 605(b), I certify that this SIP revision will not have a significant economic impact on a substantial number of small entities. (See 46 FR 8709.)

The Office of Management and Budget has exempted this rule from the requirements of section 3 of Executive Order 12291.

List of Subjects in 40 CFR Part 52

Air pollution control, Intergovernmental relations. Date: August 25, 1987.

A. James Barnes,

Acting Administrator.

Subpart L, Part 52 of Chapter I, Title 40, of the Code of Federal Regulations, is amended as follows:

PART 52-[AMENDED]

Subpart L-Georgia

1. The authority citation for Part 52 continues to read as follows:

Authority: 42 U.S.C. 7401-7642.

2. Section 52.581 is amended by redesignating paragraph (c) as paragraph (a) and by adding a new paragraph (b) to read as follows:

§ 52.581 Significant deterioration of air quality.

(b) A letter of commitment concerning the incorporation of EPA's revised modeling guidelines for PSD into the Georgia regulations was submitted to EPA on May 11, 1987, by the Georgia Department of Natural Resources.

[FR Doc. 87-19914 Filed 8-31-87; 8:45 am] BILLING CODE 6560-50-M

40 CFR Part 52

[A-1-FRL-3253-9]

Approval and Promulgation of Implementation Plans; Rhode Island; Adoption of EPA Approved Test Methods

AGENCY: Environmental Protection Agency (EPA)

ACTION: Final rulemaking.

SUMMARY: EPA is approving State Implementation Plan (SIP) revisions submitted by the State of Rhode Island. These revisions specify EPA-approved test methods to be used for compliance determinations with Rhode Island Air Pollution Control Regulations No. 11, No. 19, and No. 21 which regulate sources of volatile organic compounds (VOCs). These SIP revisions are necessary because Regulation No. 11 does not specify a test method to be used to determine compliance, and Regulations No. 19 and No. 21 require test methods which are outdated. The intended effect of this action is to include current EPAapproved testing methods in these Rhode Island Air Pollution Control Regulations of the SIP. This action is being taken under section 110 of the Clean Air Act.

effective DATE: This action will be effective on November 2, 1987, unless notice is received by October 1, 1987, that adverse or critical comments will be submitted.

ADDRESSES: Comments may be mailed to Louis F. Gitto, Director, Air Management Division, Room 2311, EPA Region I, JFK Federal Building, Boston, MA 02203. Copies of the submittal and EPA's evaluation are available for public inspection during normal business hours at the Environmental Protection Agency, Room 2311, EPA Region I, JFK Federal Building, Boston, MA 02203; Public Information Reference Unit, Environmental Protection Agency. 401 M Street, SW., Washington, D.C. 20460; and the Department of Environmental Management, 75 Davis Street, Cannon Building, Room 204, Providence, RI 02908.

FOR FURTHER INFORMATION CONTACT: Robert C. Judge, (617) 565-3248; FTS 835-3248.

SUPPLEMENTARY INFORMATION: On February 27, 1987, the Rhode Island Department of Environmental Management (DEM) submitted revisions to the Rhode Island SIP. These revisions specify EPA-approved test methods to be used for compliance determinations with Rhode Island Air Pollution Control Regulation Nos. 11, 19, and 21 which regulate VOC sources. For Regulation No. 11, the DEM is including a method for testing of leaks from bulk gasoline plants. Previously, that regulation contained no such method. For Regulations No. 19 and No. 21, which deal with volatile organic compound emissions from surface coating and printing operations, respectively, the DEM is replacing the test methods previously specified for use in compliance determinations with the test methods that are currently recommended by EPA.

For Regulation No. 11, subsection 11.4.5, the DEM is requiring the use of the test methods in Appendixes B and C of EPA publication Number 450/2–78–051 entitled Control of Volatile Organic Compound Leaks from Gasoline Tank Trucks and Vapor Collection Systems.

For Regulation No. 19, subsection 19.7.1, and Regulation No. 21, subsection 21.6.1, the DEM is requiring that Methods 24, 24A, and 25 found in Appendix A of 40 CFR Part 60 be used for compliance determinations. The amendments to Regulations No. 19 and No. 21 are written in such a way as to allow modification of the method or the use of another method as long as they are accepted by the DEM and approved by EPA. These methods are consistent with the recommended methods of the September 14, 1984 memorandum entitled "Volatile Organic Compound (VOC) Test Methods or Procedures for Source Categories in Groups I, II, and III Control Techniques Guidelines (CTGs)."

Final Action

EPA is approving revisions to Rhode Island SIP Regulations Nos. 11, 19, and 21 which specify the current EPA-approved test methods to be used to determine compliance with the requirements of these regulations.

EPA is approving these SIP revisions without prior proposal because the Agency views them as noncontroversial amendments and anticipates no adverse comments. This action will be effective 60 days from the date of this Federal Register unless, within 30 days of its publication, notice is received that adverse or critical comments will be submitted. If such notice is received, this action will be withdrawn before the effective date by publishing two subsequent notices. One notice will withdraw the final action and another will begin a new rulemaking by announcing a proposal of the action and establishing a comment period. If no such comments are received, the public is advised that this action will be effective November 2, 1987.

Under 5 U.S.C. 605(b), I certify that this SIP revision will not have a significant economic impact on a substantial number of small entities (see 46 FR 8709).

The Office of Management and Budget has exempted this rule from the requirements of section 3 of Executive Order 12291.

Under section 307(b)(1) of the Act, petitions for judicial review of this action must be filed in the United States Court of Appeals for the appropriate circuit by November 2, 1987. This action may not be challenged later in proceedings to enforce its requirements (See 307 section (b)(2).)

List of Subjects in 40 CFR Part 52

Air pollution control, Hydrocarbons, Incorporation by reference, Intergovernmental relations, Ozone, Reporting and recordkeeping requirements.

Note.—Incorporation by reference of the State Implementation Plan for the State of Rhode Island was approved by the Director of the Federal Register on July 1, 1982.

Date: August 21, 1987.

A. James Barnes,

Acting Administrator.

Subpart 00, Part 52 of Chapter I, Title 40 of the Code of Federal Regulations is amended as follows:

PART 52—[AMENDED] Subpart 00—Rhode Island

1. The authority citation for Part 52 continues to read as follows:

Authority: 42 U.S.C. 7401-7642.

2. Section 52.2070, is amended by adding paragraph (c)(30) to read as follows:

§ 52.2070 Identification of plan.

(c) * * *

(30) Revisions to the State
Implementation Plan were submitted by
Rhode Island Department of
Environmental Management on
February 27, 1987. These revisions were
effective as of January 20, 1987 in the
State of Rhode Island.

(i) Incorporation by reference. (a)
Letter from the Rhode Island
Department of Environmental
Management dated February 27, 1987
submitting revisions to the Rhode Island
State Implementation Plan.

(b) Amendment to Air Pollution Control Regulation No. 11, at subsection 11.4.5 adopted on January 20, 1987 in Rhode Island.

(c) Amendment to Air Pollution Control Regulation No. 19, at subsection 19.7.1 adopted on January 20, 1987 in Rhode Island.

(d) Amendment to Air Pollution Control Regulation No. 21, at subsection 21.6.1 adopted on January 20, 1987 in Rhode Island.

3. In §52.2081, the Table 52.2081 is amended by adding the following entrees. The date approved by EPA and the Federal Register Citation will be the publication date and citation of today's document.

§ 52.2081 EPA-Approved EPA Rhode Island State regulations.

Table 52:2081—EPA-APPROVED RULES AND REGULATIONS.

State citation	Title/subject	Date adopted by State	Date approved by EPA	FR citation	52.2070	Comments/unapproved sections
No. 11.	Petroleum liquids marketing and storage	1/20/87	9/1/87	52 FR	(c)(30)	Amended Regulation No. 11, Subsec
	Control of VOCs from Surface Coating Operations.	1/20/87	9/1/87	52 FR	(c)(30)	tion 11.4.5. Amended Regulation No. 19, Subsection 19.7.1.
No. 21	Control of VOCs from Printing Operations	1/20/87	9/1/87	52 FR	(c)(30)	Amended Regulation No. 21, Subsection 21.6.1.

[FR Doc. 87-19772 Filed 8-31-87; 8:45 am] BILLING CODE 6560-50-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

42 CFR Part 413

[BERC-435-F]

Medicare Program; Changes to the Return on Equity Capital Provisions for Outpatient Hospital Services

AGENCY: Health Care Financing Administration (HCFA), HHS. ACTION: Final rule.

SUMMARY: We are eliminating the allowance for a return on equity capital for outpatient services furnished by proprietary hospitals because it is inappropriate to continue payment of return on equity capital for outpatient hospital services while the payment for inpatient hospital services is being phased out.

EFFECTIVE DATE: For cost reporting periods beginning on or after October 1, 1987.

FOR FURTHER INFORMATION CONTACT: Anthony Coates (301) 597–2886.

SUPPLEMENTARY INFORMATION:

I. Background

Section 1861(v) of the Social Security Act (the Act) defines "reasonable cost" for Medicare purposes and provides that the necessary costs incurred by a provider (both direct and indirect) in the delivery of covered health care services are included in this definition. Currently, a return on equity capital is paid as an allowance in addition to the reasonable cost of covered services furnished to beneficiaries by proprietary skilled nursing facilities (SNFs) and proprietary hospitals, although the return on equity capital for inpatient hospital services is being phased out under section 1886(g)(2) of the Act. This section of the Act, as amended by section 9107(a) of the Consolidated Omnibus Budget Reconciliation Act of 1985 (Pub. L. 99-272) (enacted on April 7, 1986) requires a

phase-down and eventual elimination of payments for return on equity capital for inpatient hospital services. (We implemented this provision in a final rule with comment period published on June 4, 1987 (52 FR 21216).)

II. Proposed Elimination of Return on Equity Capital for Outpatient Hospital Services

On June 5, 1987, we proposed to amend 42 CFR 413.157 (at 52 FR 21330) to eliminate the allowance for a return on equity capital from payment for outpatient hospital services. In doing so, we reasoned that the allowance is unnecessary to maintain the availability of hospital outpatient services. In addition, in view of the statutory phaseout of the equity capital allowance for inpatient services as provided for under section 1886(g)(2) of the Act, we stated that continuation of the allowance for hospital outpatient services is inappropriate. We proposed that the elimination be effective for cost reporting periods beginning on or after October 1, 1987.

In that hospital capital supports inpatient and outpatient services, we stated in the proposed rule that it is inappropriate to continue payment of return on equity capital for outpatient hospital services while the payment for inpatient hospital services is being phased out. We further stated that investors in hospitals should not receive return on equity payments based upon the setting in which the hospital services are provided (that is, inpatient versus outpatient) and that the level of other Medicare payments for hospital outpatient services is adequate to maintain the availability of services to beneficiaries without an allowance for return on equity capital. In addition, hospital operating margins under the prospective payment system should encourage private investment even when payments are reduced through the elimination of return on equity capital for outpatient services. Accordingly, we concluded that the elimination of return on equity capital for outpatient hospital services is appropriate.

III. Public Comments and Response

During the public comment period, we received only one item of correspondence concerning the proposed rule. The comment was submitted by a certified public accountant and included a lengthy paper that did not deal specifically with the proposed elimination of the return on equity capital for hospital outpatient services. We have considered this commenter's arguments, but we have decided not to make any changes to the proposed rule.

Comment: The commenter stated that the allowance for a return on equity capital should not be considered to stand alone; instead, the allowance should be thought of as one part of the overall Medicare payment mechanism. The commenter maintained that eliminating the allowance for a return on equity capital as an allowable cost for the outpatient portion of proprietary hospitals would be a long term mistake. The commenter stated that the proposed rule would perpetuate a problem he believes to exist; that is, that hospitals have "no incentives to seek efficient financing arrangements" and also that the proposal might tend to discourage equity build-up. He stated that eliminating the allowance for a return on equity capital would remove yet another incentive available to encourage cost containment.

Response: Return on equity capital has been paid as an allowance in addition to the reasonable cost of covered services furnished to beneficiaries by proprietary providers. For Federal fiscal year (FY) 1988, the projected reduction in payments caused by elimination of the allowance for return on equity capital for outpatient services will represent about one-fifth of one percent of total Medicare payments to proprietary hospitals. Accordingly, the termination of payment of return on equity capital for proprietary hospital outpatient services will not adversely affect either the provider operations or beneficiary access to care.

With regard to the loss of incentive to seek efficient financing, we do not believe that the elimination of return on equity capital for hospital outpatient services creates an incentive to exercise less prudence in securing loans. When the provider finds it necessary and proper to borrow money, the Medicare program recognizes interest paid on provider debts, subject to wellestablished policies.

We also do not believe that the elimination of a return on equity capital for hospital outpatient services would tend to discourage equity build-up, especially since the return one quity represents only about one-fifth of one percent of total Medicare payments to proprietary hospitals. In fact, elimination of the return on equity capital places proprietary and non-proprietary hospitals on a more equal basis for Medicare payment purposes.

We believe that the elimination of Medicare payment of a return on equity capital may serve as an incentive to contain costs. Whereas payment for a return on equity capital used to provide an added source of income, its removal creates an added incentive to increase operational efficiency.

IV. Public Comments on the June 1987 Final Rule

In the final rule published on June 4. 1987, which we noted above, we described the phase-out of Medicare payments for a return on equity capital for inpatient hospital services that is mandated by section 1886(g)(2) of the Act, as amended by section 9107(a) of the Consolidated Omnibus Budget Reconciliation Act of 1985 (Pub. L. 99-272). We also stated, however, that under section 9321(c) of the Omnibus Budget Reconciliation Act of 1986 (Pub. L. 99-599) we were precluded from issuing final rules dealing with capitalrelated costs for inpatient hospital services until September 1, 1987. Consequently, we were unable to describe in regulations text the percent reductions imposed by section 1886(g)(2) of the Act. Therefore, in this final rule, we are adding § 413.157(b)(2) to conform the regulations to the mandate of the law.

We also want to note that, in the June 4, 1987 final rule, we provided for a 60-day public comment period on the provision that reduces the return on equity capital for all proprietary providers other than hospitals and SNFs for cost reporting periods beginning on or after October 1, 1985 (as directed by section 1861 (v)(1)(P) of the Act, as enacted by section 9107(b)(1) of Pub. L. 99-272) and before July 6, 1987 (the effective date of that final rule). We stated that we would consider all timely comments about this provision, and that if further rulemaking was necessary, we

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would publish a final rule and respond to comments in that final rule. However, we received no comments concerning this matter. Therefore, further rulemaking is unnecessary.

V. Regulatory Impact Statement

Executive Order 12291 requires us to prepare and publish a final regulatory impact analysis for regulations that are likely to have an annual effect on the economy of \$100 million or more; cause a major increase in costs or prices, or result in significant adverse effects on competition, employment, investment, productivity, innovation, or the ability of United States-based enterprises to compete with foreign-based enterprises in domestic or export markets. In addition, we generally prepare a final regulatory flexibility analysis that is consistent with the Regulatory Flexibility Act (RFA) (5 U.S.C. 601 through 612), unless the Secretary certifies that a final regulation will not have a significant economic impact on a substantial number of small entities. For purposes of the RFA, we treat all hospitals as small entities.

Currently, approximately sixteen percent of all hospitals participating in Medicare are classified as proprietary hospitals. Although a significant number of hospitals will be affected by this rule, we do not believe that this rule will have a substantial economic impact on individual hospitals. We expect that the reduction in payments to proprietary hospitals for outpatient services will range from \$15 million in FY 1988 to about \$45 million in FY 1992. While some hospitals may be more severely affected than others, we estimate that the average reduction in Medicare revenue for each proprietary hospital will represent about two percent of total payments to proprietary hospitals for outpatient services. We generally do not regard a change in payments of less than five percent as having a substantial impact on the target population.

We do not expect that beneficiary access will be affected, since most hospital outpatient departments are not located in proprietary settings. Furthermore, we believe that any incentives to limit access by proprietary outpatient departments will be more than countered by the more generalized pressures on hospitals to increase the availability of outpatient services in order to maximize the productive capability of their facilities and to provide care in the most cost-effective setting. For example, both the use of outpatient surgical services by Medicare beneficiaries and payments for these services have increased more since the

start of the prospective payment system than before.

For these reasons, we have determined that a final regulatory impact analysis is not required for this rule. In addition, we have determined, and the Secretary certifies, that this rule will not result in a significant economic impact on a substantial number of small entities. Therefore, we have not prepared a regulatory flexibility analysis.

Also, we do not expect this rule to produce an impact that will exceed the limit for reductions in payment to hospitals or physicians established by section 9321(d) of the Omnibus Budget Reconciliation Act of 1986 (Pub. L. 99–509). That provision prohibits the Secretary from issuing any final rule or notice between October 21, 1986 and September 1, 1987 that will result in a \$50 million or greater reduction in payments to hospitals or physicians in FY 1988.

VI. Other Required Information

Paperwork Reduction Act

These changes will not impose information collection requirements; consequently, they need not be reviewed by the Executive Office of Management and Budget under the authority of the Paperwork Reduction Act of 1980 (44 U.S.C. 3501 through 3511).

List of Subjects in 42 CFR Part 413

Administrative practice and procedure, Health facilities, Health professions, Kidney diseases, Laboratories, Medicare, Nursing homes, Reporting and recordkeeping requirements, Rural areas, X-rays.

Title 42 CFR Part 413 is amended as set forth below:

PART 413—PRINCIPLES OF REASONABLE COST REIMBURSEMENT; PAYMENT FOR END-STAGE RENAL DISEASE SERVICES

A. The authority citation for Part 413 continues to read as follows:

Authority: Secs. 1102, 1122, 1814(b), 1815, 1833(a), 1861(v), 1871, 1881, and 1886 of the Social Security Act as amended (42 U.S.C. 1302, 1320a-1, 1395f(b), 1395g, 1395l(a), 1395x(v), 1395hh, 1395rr, and 1395ww).

B. Section 413.157 is amended by adding new paragraph (b)(2) (previously reserved), revising paragraph (b)(3), redesignating paragraph (b)(4) as new paragraph (b)(5), and adding a new paragraph (b)(4), to read as follows:

§ 413.157 Return on equity capital of proprietary providers.

- (b) General rule. * * *
- (2) Rate of return for inpatient hospital services furnished by proprietary hospitals. The rate used in determining the return for inpatient hospital services is a percentage of the average of the rates of interest described in paragraph (b)(1) of this section. The percentages applicable to inpatient hospital services are as follows:
- (i) 150 percent for cost reporting periods beginning before April 20, 1983.
- (ii) 100 percent for cost reporting periods beginning on or after April 20, 1983 and before October 1, 1986.
- (iii) 75 percent for cost reporting periods beginning on or after October 1, 1986 and before October 1, 1987.
- (iv) 50 percent for cost reporting periods beginning on or after October 1, 1987 and before October 1, 1988.
- (v) 25 percent for cost reporting periods beginning on or after October 1, 1988 and before October 1, 1989.
- (vi) Zero percent for cost reporting periods beginning on or after October 1, 1989.
- (3) Rate of return related to proprietary SNFs. For cost reporting periods beginning on or after October 1, 1985, the rate used in determining the return for SNF's is a percentage equal to the average of the rates of interest described in paragraph (b)(1) of this section.
- (4) Rate of return related to outpatient hospital services. (i) For cost reporting periods beginning on or after October 1, 1985 but before October 1, 1987, the rate used in determining the return for outpatient hospital services is a percentage equal to the average of the rates of interest described in paragraph (b)(1) of this section.
- (ii) For cost reporting periods beginning on or after October 1, 1987, there is no allowance for return for outpatient hospital services.

(Catalog of Federal Domestic Assistance Program No. 13.773, Medicare—Hospital Insurance: and No. 13.774, Medicare— Supplementary Medical Insurance)

Dated: August 20, 1987.

William L. Roper,

Administrator, Health Care Financing Administration.

Approved: August 24, 1987.

Otis R. Bowen,

Secretary.

[FR Doc. 87-19987 Filed 8-27-87; 12:15 pm]

FEDERAL COMMUNICATIONS COMMISSION

47 CFR Parts 36 and 67

[CC Docket Nos. 78-72, 80-286 and 86-297; 87-272]

Common Carrier Services; MTS and WATS Market Structure; Amendment of the Commission's Rules and Establishment of a Joint Board

AGENCY: Federal Communications Commission.

ACTION: Final rule.

SUMMARY: The Federal Communications Commission grants, in part, reconsideration of its decision to adopt revisions of the Separations Manual, new Part 36 of its rules, recently recommended by the Federal-State Joint Board. The Commission decided to reconsider its decision to exclude access revenues from the allocation factor for marketing expenses. Thus, in the new § 36.372 of the Commission's rules (effective January 1, 1988) marketing expenses will be allocated on the basis of current billings. The Commission did not act, at this time on other reconsideration issues. The Commission has also issued a Supplemental Notice of Proposed Rulemaking, summarized elsewhere in this volume, seeking comment and data on the appropriate allocation method and recovery mechanism for marketing expenses.

EFFECTIVE DATE: January 1, 1988.

ADDRESS: Federal Communications Commission, Washington, DC 20554.

FOR FURTHER INFORMATION CONTACT: Cindy Schonhaut, Special Counsel Federal-State Joint Board Matters, Accounting and Audits Division, Common Carrier Bureau, at (202) 632–7500.

SUPPLEMENTARY INFORMATION: This is a summary of the Commission's Memorandum Opinion and Order, CC Docket Nos. 78–72, 80–286 and 86–297, FCC 87–272, adopted August 14, 1987 and released August 18, 1987.

The full text of Commission decisions are available for inspection and copying during normal business hours in the FCC Dockets Branch (Room 230), 1919 M Street, NW., Washington, DC. The complete text of this decision may also be purchased from the Commission's copy contractors, International Transcription Service, 2100 M Street, NW., Suite 140, Washington, DC 20037, (202) 857–3800.

Summary of Memorandum Opinion and Order on Reconsideration

- 1. The Joint Board in CC Docket No. 86–297 recommended that the Commission adopt a new separations manual intended to conform separations procedures to the recently revised Uniform System of Accounts (USOA) and to simplify the separations process. The Commission adopted the new Separations Manual in April 1987 which will be codified as the new Part 36 of its rules and which will become effective January 1, 1988.²
- 2. Under the current separations procedures of Part 67 of the Commission's rules, the advertising and sales expenses in the current USOA Accounts 642 and 643 are allocated between the jurisdictions on the basis of current billing for local and toll services (excluding certain billings for nonaffiliated companies and those in connection with intercompany settlements.) 3 Those expenses will be included in the new Account 6610 in the revised USOA. Under new separations procedures recommended by the Joint Board and adopted by the Commission, billings for access charges will be excluded from the allocation factor for Account 6610 marketing expenses.

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3. Several parties, including the National Association of Regulatory Utility Commissioners and the state commissioners on the Joint Board at the time the Recommended Decision and Order was adopted, urged the Commission to reconsider its decision to exclude access revenues from the allocation factor for marketing for the following reasons: (1) The Joint Board and the Commission incorrectly assumed that local exchange carriers do not actively market access services; (2) the exclusion of access revenues from the allocation factor for marketing expenses will cause a shift of \$475 million in revenue requirements to the state jurisdiction; (3) this shift in revenue requirements contravenes the stated goal of the Joint Board and the Commission to minimize the revenue requirement impact of the new Separations Manual; and, (4) the Joint Board and the Commission did not

¹ Amendment of Part 67 (New Part 36) of the Commission's Rules and Establishment of a Federal-State Joint Board, CC Docket No. 88-297, FCC 87]-4, released April 8, 1987, 2 FCC Rcd 2,582 (1987) (Recommended Decision and Order).

² MTS and WATS Market Structure, Amendment of Part 67 (New Part 36) of the Commission's Rules and Establishment of Federal-State Joint Board, CC Docket Nos. 78–72, 80–286 and 86–297, FCC 87–134, released May 1, 1987, 2 FCC Rcd 2,639 (1987) (Report and Order).

^{3 47} CFR 67.363.

provide parties with an opportunity to comment on the proposed revision of the current separations rules, thus violating the Administrative Procedure Act and applicable case law.

4. The Commission decided to reconsider its decision to exclude access revenues from the allocation factor for marketing expenses. As an interim measure, the Commission adopted a new § 36.372 of its rules which will allocate Account 6610 marketing expenses on the basis of current billings. The Commission referred a permanent resolution of this issue to the Joint Board in CC Docket No. 80-286. The Commission specified certain question for comment and requested data regarding the allocation of marketing expenses. The Commission did not address, at this time, other issues raised in the petitions for reconsideration.

Regulatory Flexibility Act

5. We certify that the Regulatory Flexibility Act * is not applicable to the rule changes we are adopting in this proceeding. In accordance with the provisions of section 605 of the Act, a copy of this certification will be sent to the Chief Counsel for Advocacy of the Small Business Administration at the time of publication of a summary of this Order and Notice in the Federal Register.

Paperwork Reduction Act

6. We have analyzed the rules adopted and proposed herein with respect to the Paperwork Reduction Act of 1980 5 and have concluded that they will not impose new or modified information collection requirements on the public. Therefore, implementation of the requirements adopted and proposed herein will not be subject to approval by the Office of Management and Budget as prescribed by the Act.

Ordering Clauses

7. Accordingly, it is ordered, That the petitions for reconsideration regarding the issue of the inclusion of access revenues in the allocation factor for marketing expense in granted to the extent indicated above.⁶

8. It is further ordered, That § 36.372 of this Commission's Rules, as set forth below, is adopted effective January 1, 1988.

List of Subjects in 47 CFR Parts 36 and 67

Communications common carrier, Telephone, Uniform System of Accounts, Reporting and recordkeeping requirements, Jurisdictional separations procedures.

Federal Communications Commission.
William J. Tricarico,
Secretary.

Part 36 of Title 47 of the Code of Federal Regulations, effective January 1, 1988, is amended as follows:

PART 36-[AMENDED]

1. The authority citation for Part 38 continues to read as follows:

Authority: 47 U.S.C. secs. 151, 154(i) and (j), 205, 221(c), 403 and 410.

Section 36.372 is amended by revising paragraph (a) to read as follows:

§ 36.372 Marketing-Account 6610.

(a) The expenses in this account are apportioned among the operations on the basis of an analysis of current billing for a representative period, excluding current billing on behalf of others and billing in connection with intercompany setttlements.

[FR Doc. 87–19922 Filed 8–31–87; 8:45 am]

47 CFR Part 76

[MM Docket No. 85-349]

Carriage of Television Broadcast Stations on Cable Television Systems; Correction

AGENCY: Federal Communications Commission.

ACTION: Final rule; correction.

SUMMARY: This document corrects the final rule published in proceeding concerning carriage of television broadcast stations on cable television systems published on May 11, 1987, 52 FR 17574.

ADDRESS: Federal Communications Commission, Washington, DC 20554.

FOR FURTHER INFORMATION CONTACT: Scott Roberts, Mass Media Bureau, (202) 632–6302.

SUPPLEMENTARY INFORMATION: This is a summary of the Commission's Erratum in MM Docket No. 85–349, released August 24, 1987. The full text of this Commission decision is available for inspection and copying during normal business hours in the FCC Dockets Branch (Room 230), 1919 M Street,

Northwest, Washington, DC. The complete text of this decision may also be purchased from the Commission's copy contractors, International Transcription Services, (202) 857–3800, 1919 M Street, NW., Room 246, Washington, DC 20554.

List of Subjects in 47 CFR Part 76

Cable television.

Part 76 of Title 47 of the Code of Federal Regulations is amended as follows:

PART 76-[AMENDED]

 The authority citation for Part 76 continues to read as follows:

Authority: 47 U.S.C. 154, 303 and 521.

2. Section 76.5 is amended by revising paragraphs (d)(1) introductory text and (d)(1)(ii) to read as follows:

§ 76.5 Definitions.

(d) Qualified station. (1) Any television broadcast station, as defined in § 76.5(b), except where such station would be considered a distant signal for copyright purposes, that with respect to a particular cable system:

(ii) If a commercial station receives an average share of total viewing hours of at least two percent and a net weekly circulation of at least five percent, as defined in § 76.5(k), in noncable households in the county served by the cable system or has been operational less than one full year. For purposes of this section, a station is considered operational as of the date it initially commences operation under program test authority. Changes in station operations, for example, upgrade of facilities, transfer or assignment of license, or recommencement after operations have ceased, are not considered initial commencement of operations under this paragraph. The viewing standards of this paragraph shall not apply for one full year from June 10, 1987, to otherwise qualified stations that commenced operation on or after July 19, 1985, but before June 10, 1987 (the effective date of these rules). Once a commercial station has demonstrated that, on the basis of a full one-year survey season, it meets the viewing standard, it will be considered to have satisfied this standard for the remainder of the period until June 10, 1982; Provided, however, that at any time after the viewing standard, a cable system may nullify the station's mandatory signal carriage eligibility if it demonstrates that it meets the viewing standard, a cable system may nullify the

⁴⁵ U.S.C. 603.

^{5 44} U.S.C. 501.

⁶ This action is taken pursuant to 47 U.S.C. 154(i) and (j). 201, 202, 205, 218, 221(c), 403 and 410.

station's mandatory signal carriage eligibility if it demonstrates, using the methodology specified in § 76.5 of this part, that the station no longer meets the viewing standard.

Federal Communications Commission.
William J. Tricarico,
Secretary.

[FR Doc. 87-19928 Filed 8-31-87; 8:45 am]

DEPARTMENT OF TRANSPORTATION

Research and Special Programs
Administration

49 CFR Part 192

[Docket No. PS-84, Amdt. 192-56]

Transportation of Natural and Other Gas by Pipeline; Confirmation or Revision of Maximum Allowable Operating Pressure Near Certain Occupied Buildings and Outside Areas

AGENCY: Research and Special Program Administration (RSPA), DOT. ACTION: Final rule.

SUMMARY: This final rule amends the criteria used to classify pipelines located near certain buildings and outside areas that are occupied infrequently. The effect is to relieve the undue burdens imposed by the current rules when pipelines are near these buildings or areas. Considering the risk, an acceptable level of safety will still be provided by the revised criteria and applicable safety standards.

EFFECTIVE DATE: October 1, 1987.

FOR FURTHER INFORMATION CONTACT: Mr. Paul J. Cory, (202) 366–4561 regarding the content of this amendment or Ms. Sandra Cureton, Dockets Unit, Office of Hazardous Materials (202) 366– 5046 regarding copies of the amendment or other information in this docket.

SUPPLEMENTARY INFORMATION:

Background

Notice 1 of this proceeding (50 FR 36116, September 5, 1985) (ANPRM) explained that this rulemaking is a result of requests from five pipeline operators for waiver of § 192.611 as it pertains to pipelines that have been reclassified according to criteria under § 192.5(d)(2). Section 192.611 requires confirmation or revision of maximum allowable operating pressure (MAOP) in areas where there has been population growth as represented by an increase in class location under the criteria of § 192.5. The waiver requests involved

pipelines built to class location 1 standards which had under gone a class location jump from 1 to 3. This normally involves replacement of the line section, although reduction in operating pressure is also a permissible remedy.

The criteria of § 192.5(d)(2) are:

§ 192.5 Class locations.

(d) A Class 3 location is:

(2) An area where the pipeline lies within 100 yards of any of the following:

 (i) A building that is occupied by 20 or more persons during normal use.

(ii) A small, well-defined outside area that is occupied by 20 or more persons during normal use, such as a playground, recreation area, outdoor theater, or other place of public assembly.

The waiver petitions cited the high costs of confirming or revising the MAOP for short segments of pipeline (approximately 600 feet each), the small number of occupants of the buildings or outside areas, and the infrequency of occupancy (such as once or twice a week) to argue that the required confirmation or revision in MAOP was not justified. The requests were not granted, however, because none of the operators demonstrated that public safety would not be adversely affected if the MAOP of the pipeline segment or segments involved were not confirmed or revised as required by § 192.611. Nevertheless, RSPA observed that § 192.5(d)(2) may be too conservative when compared to other class location criteria, and some softening of the criteria might be accomplished without a reduction in safety

In the ANPRM RSPA requested comments on six alternative that were seen as possible courses of action, and asked eight questions relating to the application of § 192.611 under the class location 3 described in § 192.5(d)(2).

An analysis of available information and comments to the ANPRM was published in Notice 3 (NPRM) (51 FR 29504, August 18, 1986). In the NPRM, RSPA proposed to amend § 192.5(d)(2) by deleting the phrase "during normal use" for both buildings and outside areas that are occupied by 20 or more persons and by replacing the deleted phrase with "on at least 5 days a week during at least 26 weeks a year." This proposal was designed to quantify the risk exposure represented by § 192.5(d)(2), and to set the level of exposure high enough that occasional usages, such as 1-week county fairs or rural churches, would not, by themselves, trigger class 3 responses under Part 192, either under § 192.611 or other rules.

Discussion of Comments to the NPRM

Twenty nine commenters responded to the NPRM. Summarized comments and RSPA responses are:

Comment #1: Fourteen comments agreed with the wording proposed in the NPRM for § 192.5(d)(2).

RSPA Response: None.

Comment #2: Six comments agreed in principle with the proposal but pointed out that for clarity the regulations should state that neither the days nor the weeks have to run consecutively. They also said that a year should be any 12-month period beginning with the date of the first known occupancy by more than 20 or more persons.

RSPA Response: For the most part.
RSPA agrees with these comments.
Since the exposure is the same, the days and weeks do not have to run consecutively and a year need not be a calendar year. Appropriate changes have been made in the final rule to make this clear. Although RSPA also agrees that the 12-month period starts with the time that 20 or more people are known to have been in occupancy, it would not be reasonable to apply the criteria otherwise, and so we see no need to specify the beginning point of the 12-month period.

Comment #3: Four comments agreed with the proposal but recommended that "20 or more persons" be increased to some larger number of persons. One of these did not mention a number, however, the other two comments recommended changing 20 to 100 persons. One commenter suggested that we consider an additional class location 3 designation that would apply where pipelines lay within 100 yards of an area where 500 or more persons assemble at least 10 days per year.

RSPA Response: The idea of raising the number of persons from 20 to some greater number was discussed as an alternative in the ANPRM. Sixty-five percent of the comments to that notice did not believe it would alleviate the problem. It would also expose more people to risk. Therefore, RSPA did not propose to change the number in the NPRM. Rather, we proposed to quantify the frequency of use, or length of exposure of 20 or more persons to the pipeline, as the best way to resolve the problem of infrequent usage, while minimizing undesired effects on safety.

The comment regarding 500 or more persons for at least 10 days a year was not adopted because the high occupancy type of usage this commenter had in mind nevertheless falls in the realm of occasional exposure to risk to which the NPRM was directed. The overall

exposure of 500 people for 10 days probably would be no greater than that of a county fair. County fairs, which involve large congregations of people for about a week, were mentioned in the NPRM as a type of occasional usage that does not deserve the more stringent Class 3 treatment. No comments were voiced in opposition to excluding county fairs from Class 3 designations.

Comment #4: One commenter agreed with the proposal but recommended that the 26 weeks per year be changed to 13 weeks because most schools observe a variety of holidays that could result in their not having 26 weeks with 5 days

per week.

RSPA Response: As a result of this comment on schools and comment #5, RSPA has reduced the occupancy period to 10 weeks, which appears sufficient to satisfy the need to protect schools as well as the objections discussed in comment #5.

Comment #5: Two commenters pointed out that 5 days per week and 26 weeks per year would exclude resort areas such as theme parks, summer camps, camp grounds, public swimming pools, etc., that would be occupied in many areas from ¼ to ½ of the year.

RSPA Response: We agree that areas or buildings such as these that are normally occupied by a large number of persons during a few months of the year warrant additional consideration. Such facilities as theme parks, summer camps, camp grounds, and public swimming facilities, etc., in most areas of the U.S., are open by the last week in June or the first week of July and remain open at least until Labor Day. This is a period of 10 weeks and may be 11 weeks. They usually are open at least 5 days a week. A few summer camps may be in session for only eight or nine weeks. OPS intends to include these summer camps within the rules protection. We have drafted the rule in terms of 10 weeks, rather than 8 or 9 because we believe that these full-length summer camps usually will have 20 or more staff and/or other persons present in the weeks before and after the weeks when the camps are in session to prepare for or shut down the camp. The reduction from 26 to 10 weeks should not affect the objective of excluding occasionally used facilities since usually they are not in session 5 days a week or for 10 weeks. As a result of this comment and comment #4, the final rule reduces the number of weeks from 26 to 10.

Comment #6: One comment recommended that the present wording of § 192.5(d)(2) be retained, but that the requirement of § 192.611 to confirm or revise the MAOP be waived for such Class 3 areas. Thus, the pressure would not have to be reduced or the pipe replaced, but all other monitoring and maintenance requirements applicable to that Class 3 location would remain.

RSPA Response: This also was one of the six alternatives mentioned in the ANPRM. Two thirds of the commentors to the ANPRM rejected the idea. RSPA did not propose it in the NPRM because of the uncertain effect on safety of excepting all § 192.5(d)(2) Class 3 locations from the requirements of § 192.611.

Comment #7: Three commenters recommended adopting the appropriate provisions from the American Society of Mechanical Engineers B31.8 Code—(1984a edition), a voluntary code of standards for gas piping systems.

RSPA Response: Although the B31.8 wording provides good guidelines for dealing with the subject conditions, this comment was not adopted because the vague B31.8 language would allow wide variations in the level of safety provided in similar locations and reduce the enforceability and effectiveness of the § 192.5(d)(2) criteria.

Comment #8: One commenter recommended no change in the present rule but a more liberal use of waivers based on the specifics of each case and the recommendations of the regulatory agency responsible for pipeline safety in

the State involved.

RSPA Response: This concept was discussed in the NPRM in response to ANPRM alternate #1. It was rejected because the problem areas are too numerous to handle on a waiver, or case by case, basis.

Advisory Committee Review

Section 4(b) of the Natural Gas Pipeline Safety Act of 1968, as amended (49 U.S.C. 1673(b)), requires that each proposed amendment to a safety standard established under this statute be submitted to a 15-member advisory committee for its consideration. The **Technical Pipeline Safety Standards** Committee, composed of persons knowledgeable about transportation of gas by pipeline discussed the proposed rule at a meeting held June 10, 1986. The Committee unanimously voted that the proposal was technically feasible, reasonable and practicable. The Committee's official report for the meeting is in the docket.

Classification

This final rule is considered to be nonmajor under Executive Order 12291 and is not a significant rule under DOT Regulatory Policies and Procedures (44 FR 11034, February 26, 1979). The economic impact of this final rule will amount to about 24 million dollars average annual savings for the industry and consumers.

Since the impact of this final rule is expected to affect primarily operators of transmission pipelines, the agency certifies that it will not have a significant economic impact on a substantial number of small entities.

List of Subjects in 49 CFR Part 192

Pipeline safety, Class location, Maximum allowable operating pressure. In view of the foregoing RSPA amends 49 CFR Part 192 as follows:

PART 192-[AMENDED]

1. The authority citation for Part 192 continues to read as follows:

Authority: 49 U.S.C. 1672; 49 U.S.C. 1804; 49 CFR 1.53 and Appendix A of Part 1.

2. In § 192.5 paragraph (d)(2) is revised to read as follows:

§ 192.5 Class locations.

(d) * * *

(2) An area where the pipeline lies within 100 yards of either a building or a small, well-defined outside area (such as a playground, recreation area, outdoor theater, or other place of public assembly) that is occupied by 20 or more persons on at least 5 days a week for 10 weeks in any 12-month period. (The days and weeks need not be consecutive.)

Issued in Washington, DC, on August 26,

M. Cynthia Douglass,

Administrator, Research and Special Programs Administration. [FR Doc. 87–19907 Filed 8–31–87; 8:45 am] BILLING CODE 4910–60–M

Federal Highway Administration

49 CFR Part 383

Commercial Driver's License Standards; Technical Correction

AGENCY: Federal Highway Administration, DOT.

ACTION: Final rule; technical correction.

SUMMARY: This document corrects a rule on commercial driver's license standards that appeared at page 20574 in the Federal Register of Monday, June 1, 1987 [52 FR 20574]. This action is necessary to correct a typographical error in § 383.37, Employer responsibilities.

EFFECTIVE DATE: September 1, 1987.

FOR FURTHER INFORMATION CONTACT:

Mr. Michael F. Trentacoste, Office of Motor Carrier Standards (202) 366–4009, or Mr. Michael J. Laska, Office of the Chief Counsel (202) 366–1383, Federal Highway Administration, 400 Seventh Street, SW., Washington, DC 20590. Office hours are from 7:45 a.m. to 4:15 p.m., e.t., Monday through Friday, except legal holidays.

§ 383.37 [Corrected]

In FR Doc. 87–12467, in the issue of Monday, June 1, 1987, on page 20588, in 49 CFR 383.37, substitute the word "employer" for the word "employee" in the introductory paragraph of the section.

(23 U.S.C. 315; 49 CFR 1.48)

Issued on: August 25, 1987.

Hugh T. O'Reilly,

Deputy Chief Counsel, Federal Highway Administration.

[FR Doc. 87-20000 Filed 8-31-87; 8:45 am] BILLING CODE 4910-22-M

DEPARTMENT OF THE INTERIOR

Fish and Wildlife Service

50 CFR Part 17

Endangered and Threatened Wildife and Plants; Final Rule to Determine Penstemon Haydenii (Blowout Penstemon) To Be an Endangered Species

AGENCY: Fish and Wildlife Service, Interior.

ACTION: Final rule.

SUMMARY: The Service determines a plant, Penstemon haydenii (blowout penstemon), to be an endangered species under the authority of the Endangered Species Act of 1973, as amended. Critical habitat is not being designated. The blowout penstemon is known from small populations in Cherry (3 populations), Hooker (1 population), Garden (3 populations), Box Butte (2 populations), and Sheridan (1 population) Counties, Nebraska. The number of plants estimated in 1986 in all populations was 2,100 ± 200. The number of plants varies considerably from year to year.

Approximately 40 to 45 percent of the populations are located on private and State lands, and 55 to 60 percent are located on Service lands. The stablization of blowout complexes leads to declining numbers of the species. The low probabilities of seed fertilization, maturation, and dispersal and seedling establishment may also contribute to the decline of the species. This

determination that *Penstemon haydenii* is endangered implements the protection provided by the Act.

DATES: The effective date of this rule is October 1, 1987.

ADDRESSES: The complete file for this rule is available for inspection, by appointment, during normal business hours at the Nebraska Field Office, U.S. Fish and Wildife Service, 2604 St. Patrick, Suite 7, Grand Island, Nebraska 68803.

FOR FURTHER INFORMATION CONTACT: Mr. Wally Jobman, Staff Biologist, Fish and Wildlife Enhancement Division, Endangered Species Office, at the above address (308/381-5571 or FTS 541-6571). SUPPLEMENTARY INFORMATION:

Background

Penstemon haydenii (blowout penstemon) was described by Sereno Watson (1891), based on a collection by H. L. Webber near Dismal River in Thomas County, Nebraska. The plant was also found there in 1889 by Webber and perhaps earlier by F.V. Hayden.

Penstemon haydenii, a member of the snapdragon family, is a hairless perennial that grows 1 to 2 feet high. The stems are often decumbent, simple or branched, and very leafy. The stem leaves are linear to lanceolate, entire, 3 to 5 inches long by 1 to 3 inches wide, sessile and clasping. The inflorescence is a compactly crowded thyrse. Floral bracts are ovate to lanceolate, nearly equalling the flowers. The corolla is blue and 1.5 to 2 inches long. Penstemon haydenii can be distinguished from P. angustifolius by its larger and lighter blue flowers. The species flowers from mid-May to late June. The flowers have a strong, persistent fragrance that lures seveal kinds of bees and other pollinators.

Historically, Penstemon haydenii probably was widely scattered throughout the central part of the Sandhills of Nebraska. All herbarium specimens and most literature citations indicate that it has never been collected outside of Nebraska. A purported Wyoming collection by Hayden was reported as being from Nebraska (Pennell 1935, p. 269), and reports of the species from Kansas are believed to be misidentifications (Craig Freeman, University of Connecticut, personal communication) and are not accepted in the Atlas of the Flora of the Great Plains (Barkley 1977).

The species is restricted to active blowouts in the sandhills of Cherry, Hooker, Box Butte, Sheridan, and Garden Counties, Nebraska, and many historic locations do not support the species today because of elimination of the habitat due to stablization of the sand dunes as a range management practice.

All know sites are well-developed blowouts in dune complexes with active sand and accompanying environmental extremes in wind, temperature, evapotranspiration, and soil moisture stress. Penstemon haydenii is found most frequently in microsites that are, or recently have been, zones of sand accumulation. The plant apparently is successional and is a primary invader that does not persist when a blowout becomes completely vegetated (Pool 1914). The species survives burial in sand by sending off shoots at successively higher nodes. It withstands initial erosion but does not have the rhizomatous system or extensive lateral roots to survive erosion that uncovers much more than a few inches of root length.

In the December 15, 1980, Federal Register (45 FR 82480), the Service published a notice of review for plants under consideration for listing as endangered or threatened, including Penstemon haydenii. A second notice of review for plants was published September 27, 1985, in the Federal Register (50 FR 39526) and included Penstemon haydenii as a category 1 species. All candidate taxa in the 1985 notice are treated as under petition (48 FR 53641).

On February 15, 1983, the Service published a notice (48 FR 6752) of its prior finding that substantial scientific information had been presented that indicates that the petitioned action on this species may be warranted in accordance with section 4(b)(3)(A) of the Endangered Species Act of 1973, as amended (the Act). On October 13, 1983, October 12, 1984, and October 11, 1985, petition findings were made that listing Penstemon haydenii was warranted but precluded by other pending listing actions, in accordance with section 4(b)(3)(B)(iii) of the Endangered Species Act. Such finding requires a recycling of the petition pursuant to a section 4(b)(3)(C)(i) of the Act. On April 29, 1986, the Service published a proposed rule (51 FR 15929) to list Penstemon haydenii as an endangered species, constituting the next required 1-year finding.

Summary of Comments and Recommendations

In the April 29, 1986, proposed rule (51 FR 15929) and associated notifications, all interested parties were requested to submit factual reports or information that might contribute to the development of a final rule. Appropriate State

agencies, county governments, Federal agencies, scientific organizations, and other interested parties were contacted and requested to comment. Newspaper notices that invited general public comment were published in the Omaha World Herald (May 29–31 and June 1) and in the Valentine Newspaper, Garden County News, and Hooker County Tribune on May 29 and on June 5, 12, and 19, 1986. Seven comments were received and are discussed below. No public hearing was requested or held.

The Ainsworth Irrigation District expressed concern that landowners will not be allowed to repair and control blowouts if this species is listed as endangered. The Service responds that private landowners are not subject to any taking prohibitions for plants listed under the Endangered Species Act. However, any Federal agency that funds, authorizes, or carries out an action in the area where the blowout penstemon may be present must ensure under section 7 of the Endangered Species Act that its action is not likely to jeopardize the continued existence of the species. Species management activities may be needed to protect and recover the species which may affect some local actions that receive Federal funds for land stabilization activities.

The Nebraska Game and Parks Commission supported the listing and agreed that designation of critical habitat would not be prudent. The Commission brought to our attention that the statement, under the Summary of Factors Affecting the Species, that Penstemon haydenii is not protected by any State laws or regulations, was in error. On January 8, 1986, the Commission took formal action and listed the blowout penstemon as an endangered species under the authority of the Nebraska Nongame and Endangered Species Conservation Act. The Commission suggested Box Butte County be added to the list of county occurrences and that only 55-60 percent of the populations occur on Federal lands. According to the Commission, justifying listing on the basis of the number of individuals may not be appropriate because of the large fluctuation in plant numbers from yearto-year. The Service has considered these corrections and suggestions and has made appropriate changes in this final rule.

A professor at the University of Nebraska, Department of Agronomy, supported the listing and submitted his estimate of 2,100 ± 200 plants as being the 1986 population. He estimated the population percentage on Federal land

to be between 25 and 50 percent. He also commented that population numbers tend to fluctuate greatly from year to year. The Service has incorporated these comments into the final listing.

Whiskey Basin Consultants commented that currently available information supports listing this species as endangered, but that additional surveys are needed to further document the occurrence of blowout penstemon. The Soil Conservation Service (SCS) questioned the advisability of listing the species without further survey of its range and pointed out that Penstemon haydenii is now listed as a State endangered species. However, no supporting biological information was received from these commenters, and the Service's decision to list the species is based on the best information currently available, which indicates a small number of populations.

A former part-owner of a site supporting Penstemon haydenii concurred that the species' numbers have decreased in the past 50 years and recommended methods to increase the blowout habitat.

Summary of Factors Affecting the Species

After a thorough review and consideration of all information available, the Service has determined that Penstemon haydenii should be classified as an endangered species. Procedures found at section 4(a)(1) of the Endangered Species Act (16 U.S.C. 1531 et seq.) and regulations (50 CFR Part 424) promulgated to implement the listing provisions of the Act were followed. A species may be determined to be an endangered or threatened species due to one or more of the five factors described in section 4(a)(1). These factors and their application to Penstemon haydenii Watson (blowout penstemon) are as follows:

A. The Present or Threatened Destruction, Modification, or Curtailment of Its Habitat or Range

Successful control of unstable sand dunes has resulted in restriction of the required blowout habitats of *Penstemon haydenii*. The blowouts where the species grow are conical or irregularly-shaped craters that are scooped out of sand by the swirling action of prevailing westerly winds. Because of successful dune stabilization programs that protect farmlands in the sandhills, the species does not have adequate habitat to invade. The decrease in extent of blowouts also has made dispersal to the fewer remaining natural blowouts more difficult.

B. Overutilization for Commercial, Recreational, Scientific, or Educational Purposes

The species is attractive and has been cultivated. Horticultural collecting is a potential threat for such a species known from so few individuals.

C. Disease or Predation

None known.

D. The Inadequacy of Existing Regulatory Mechanisms

Penstemon haydenii is listed as endangered under the Nebraska Nongame and Endangered Species Conservation Act (sections 37-430 to 37-438, Nebraska Revised Statutes), which regulates possession, transportation, exportation from the State, processing, sale or offer for sale, or shipment of the species within the State. Under the provisions of 50 CFR Parts 25 through 28, the Service provides some protection for the species on refuge lands. Approximately 55-60 percent of known populations are on Service refuge land and 40-45 percent are on State and private lands. The Endangered Species Act will provide additional protection of this species through section 7 (interagency cooperation) requirements and through section 9, which, among other things, prohibits removal and reduction to possession of listed plants on areas under Federal jurisdiction.

E. Other Natural or Manmade Factors Affecting its Continued Existence

Penstemon haydenii comprises nine small populations that consist of a total of approximately 2,100 individuals. The small population size makes the species vulnerable to localized environmental changes. In addition, the species occupies a successional niche in the development and eventual revegetation of blowout habitats. As the vegetational cover in these areas increases, P. haydenii undergoes local extirpation. The species is not only rare, but does not appear vigorous at the known localities, possibly because these blowouts have reached a stage of revegetation that exceeds the optimum habitat conditions for the species, and the number of new blowouts is decreasing.

The Service has carefully assessed the best scientific and commercial information available regarding the past, present, and future threats faced by this species in determining to make this rule final. Based on this evaluation, the preferred action is to list Penstemon haydenii as endangered. With only about 2,100 individuals known and stabilization of blowout complexes

causing further declines, endangered status seems an accurate assessment of the species' condition. For the reasons stated below, no critical habitat designation is included in this rule.

Critical Habitat

Section 4(a)(3) of the Endangered Species Act, as amended, requires that, to the maximum extent prudent and determinable, the Secretary designate critical habitat at the time a species is determined to be endangered or threatened. The Service finds that designation of critical habitat is not prudent for Penstemon haydenii at this time. This species depends on early successional stages in the revegetation of sandhill blowouts for its habitat. Such blowouts are transient features of the sandhill topography, and a critical habitat designation reflecting the present habitat occupied by the species would quickly become inappropriate as present blowouts become stabilized and new ones develop. Even supposing that critical habitat could be kept in a state of revision to reflect the varying range of the species, such public identification of habitat would be inadvisable for such an attractive and conspicuous flowering plant, which could easily be exposed to vandalism or horticultural collecting. All involved parties and landowners will be notified of the location and importance of protecting this species' habitat. Protection of this species' habitat will be addressed through the recovery process and through the section 7 jeopardy standard. Thus, the Service concludes that designation of critical habitat for this species would be neither practical nor beneficial to its conservation and therefore is not prudent.

Available Conservation Measures

Conservation measures provided to species listed as endangered or threatened under the Endangered Species Act include recognition, recovery actions, requirements for Federal protection, and prohibitions against certain practices. Recognition through listing encourages and results in conservation actions by Federal, State, and private agencies, groups, and individuals. The Endangered Species Act provided for possible land acquisition and cooperation with the States and requires that recovery actions be carried out for all listed species. Such actions are initiated by the Service following listing. The protection required of Federal agencies and the prohibitions against collecting and trade are discussed, in part, below.

Section 7(a) of the Act, as amended. requires Federal agencies to evaluate their actions with respect to any species that is proposed or listed as endangered or threatened and with respect to its critical habitat if it is being designated. Regulations implementing this interagency cooperation provision of the Act are codified at 50 CFR Part 402. Section 7(a)(2) requires Federal agencies to ensure that activities they authorize, fund, or carry out are not likely to jeopardize the continued existence of a listed species or to destroy or adversely modify its critical habitat. If a Federal action may affect a listed species or its critical habitat, the responsible Federal agency must enter into formal consultation with the Service. Some management actions, such as stabilization of sand dunes by the U.S. Fish and Wildlife Service and the Soil Conservation Service, might adversely impact this species, since stabilization deprives the plant of suitable habitat for growth and reproduction. There may be a need for consultation under section 7 regarding the Soil Conservation Service's partial funding of private erosion-control activities. The Fish and Wildlife Service will be responsible for assuring that management of the two National Wildlife Refuges on which this species occurs is consistent with maintaining its continued survival. The Service will also seek voluntary cooperation with private landowners in managing habitat suitable for this species, and may undertake reestablishment of populations within former range on Federal or other lands.

The Endangered Species Act and its implementing regulations found at 50 CFR 17.61 and 17.62 set forth a series of general prohibitions and exceptions that apply to all endangered plants. All prohibitions of section 9(a)(2) of the Act, implemented by 50 CFR 17.61, apply. These prohibitions, in part, make it illegal for any person subject to the jurisdiction of the United States to import or export any endangered plant, transport it in interstate or foreign commerce in the course of a commercial activity, sell or offer it for sale in interstate or foreign commerce, or remove it from areas under Federal jurisdiction and reduce it to possession. Certain exceptions can apply to agents of the Service and State conservation agencies. The Act and 50 CFR 17.62 also provide for the issuance of permits to carry out otherwise prohibited activities involving endangered plant species under certain circumstances. With

respect to *P. haydenii*, few permits are expected to be sought or issued, since the species is not common in cultivation or in the wild. Requests for copies of the regulations on plants and inquiries regarding them may be addressed to the Federal Wildlife Permit Office, U.S. Fish and Wildlife Service, Washington, DC 20240 (703/235–1903).

National Environmental Policy Act

The Fish and Wildlife Service has determined that an Environmental Assessment, as defined under the authority of the National Environmental Policy Act of 1969, need not be prepared in connection with regulations adopted pursuant to section 4(a) of the Endangered Species Act of 1973, as amended. A notice outlining the Service's reasons for this determination was published in the Federal Register on October 25, 1983 (48 FR 49244).

References Cited

Barkley, T.M., editor. 1977. Atlas of the Flora of the Great Plains. The Iowa State University Press. 578 pages.

Pennell, F.W. 1935. Scrophulariaceae of East Temperate North America. Academy of Natural Sciences of Philadelphia Monographs, 1:267–269.

Pool, R.J. 1914. A study of the vegetation of the Sandhills of Nebraska, Minnesota Botanical Studies, 3(4):189.

Smyth, B. 1899. Additions to the Flora of Kansas. Proceedings and Transactions of the Kansas Academy of Science, 6:158–167. Watson, S. 1891. Penstemon haydenii, n. sp. Botanical Gazette, 16:311.

Author

The primary author of this final rule is Mr. Wally Jobman, Nebraska Field Office, U.S. Fish and Wildlife Service, 2604 St. Patrick, Suite 7, Grand Island, Nebraska 68803. A status report was prepared by Mr. Robert W. Lichvar of Whiskey Basin Consultants, Cheyenne, Wyoming. Dr. James L. Miller of the Denver Regional Endangered Species Office served as editor.

List of Subjects in 50 CFR Part 17

Endangered and threatened wildlife, Fish, Marine mammals, Plants (agriculture).

Regulation Promulgation

Accordingly, Part 17, Subchapter B of Chapter I, Title 50 of the Code of Federal Regulations, is amended as set forth below:

PART 17-[AMENDED]

1. The authority citation for Part 17 continues to read as follows:

Authority: Pub. L. 93–205, 87 Stat. 884; Pub. L. 94–359, 90 Stat. 911; Pub. L. 95–632, 92 Stat. 3751; Pub. L. 96–159, 93 Stat. 1225; Pub. L. 97–304, 96 Stat. 1411 (16 U.S.C. 1531 et seq.).

2. Amend § 17.12(h) by adding the following, in alphabetical order under Scrophulariaceae, to the List of Endangered and Threatened Plants:

Species		tea a san	0	148-1-1-1	Critical	Special
Scientific name	Common name	Historic range	Status	When listed	Critical habitat	Special rules
	Table Street Street	The second second				
ophulariaceae—Snapdragon family: stemon haydenii	Blowout penstemon	1101 05	The state of the state of		NA	

Dated: August 3, 1987.

Susan Recce,

Acting Assistant Secretary for Fish and Wildlife and Parks.

[FR Doc. 87-20021 Filed 8-31-87; 8:45 am]
BILLING CODE 4310-55-M

Proposed Rules

Federal Register

Vol. 52, No. 169

Tuesday, September 1, 1987

This section of the FEDERAL REGISTER contains notices to the public of the proposed issuance of rules and regulations. The purpose of these notices is to give interested persons an opportunity to participate in the rule making prior to the adoption of the final rules.

DEPARTMENT OF AGRICULTURE

Food and Nutrition Service

7 CFR Part 210

Revision of Contract Duration for Food Service Management Company

AGENCY: Food and Nurition Service, USDA.

ACTION: Proposed rule.

SUMMARY: This document proposed to amend Part 210, National School Lunch Program, to increase the number of allowable annual renewals of contracts between school food authorities and food service management companies from 2 to 4. This proposal responds to a number of comments on this matter which were received by the Department in connection with the interim rule that revised and reorganized Part 210 (51 FR 34864, September 30, 1986). Since the proposal that preceded the interim rule contained no change to existing contract duration requirements, the Department wishes to provide more specific notice of its intent to address that issue than was provided in that particular rulemaking. The Department is proposing this rule to provide a potentially more stable environment for school food service operations that are conducted in whole or in part under contracts with food service management companies and to decrease the paperwork and administrative burden of school food authorities that have such contracts.

DATE: To be assured of consideration. comments must be postmarked no later than November 2, 1987.

ADDRESS: Comments should be sent to Lou Pastura, Branch Chief, Policy and Program Development Branch, Child Nutrition Division, Food and Nutrition Service, USDA, Alexandria, Virginia 22302. All written submissions will be available for public inspection in Room 509, 3101 Park Center Drive, Alexandria, Virginia 22302, during regular business

hours (8:30 a.m. to 5:00 p.m.) Monday through Friday.

FOR FURTHER INFORMATION CONTACT: Mr. Pastura at the address listed above or call (703) 756-3620.

SUPPLEMENTARY INFORMATION:

Classification

This proposed action has been reviewed under Executive Order 12291 and has been classified not major. We anticipate that this proposal will not have an impact on the economy of more than \$100 million. No major increase in costs or prices for program participants, individual industries, Federal, State or local government agencies, or geographic regions is anticipated. The proposal is not expected to have significant adverse effects on competition, employment, investment, productivity, innovation, or on the ability of U.S.-based enterprises to compete with foreign-based enterprises in domestic or export markets.

The National School Lunch Program is listed in the catalog of Federal Domestic Assistance under No. 10.555 and is subject to the provisions of Executive Order 12372 which requires intergovernmental consultation with State and local officials (See 7 CFR Part 3015, Subpart V, 48 FR 29112, June 24,

This proposal has also been reviewed with regard to the requirements of Pub. L. 96-354, the Regulatory Flexibility Act. The Acting Administrator of FNS has certified that this proposal will not have a significant economic impact on a substantial number of small entities.

This proposed action would impose no new reporting or recordkeeping provisions that are subject to Office of Management and Budget review in accordance with the Paperwork Reduction Act of 1980 (44 U.S.C. 3501 through 3520).

Background

Prior to 1970, school food authorities were prohibited from contracting with food service management companies to provide meals under the National School Lunch Program (NSLP). In 1970, the NSLP regulations were amended to authorize any school food authority to contract with a food service management company to conduct all or part of its feeding operation under the Program. No restrictions were imposed on the duration of such contracts.

The NSLP regulations were again amended in 1978 to address food service management companies. Under that amendment, additional requirements and controls were imposed upon school food authorities contracting with food service management companies. These provisions have remained essentially unchanged to date (7 CFR 210.16). Among the requirements imposed by the amendment was a contract duration period of 1 year with option for 2 renewals of 1 year each. As a result, the bidding cycle for food service management company contracts cannot exceed 3 years. In the 1978 rulemaking, the Department originally proposed that contracts be for 1 year only with no option for renewal. However, in response to substantial commenter concern over such a restrictive contract period, the Department provided the renewal option in the final rule.

On February 12, 1985, the Department published a proposed rewrite of Part 210 in the Federal Register (50 FR 5950). This proposal was intended to reorganize and clarify the NSLP regulations as well as to make several substantive policy changes. No change was proposed in the duration of food service management company contracts and only one comment was received on this provision (7 CFR 210.16(d). Consequently, no change to the provision was made in the interim rewrite of Part 210 which was published on September 30, 1986 (51 FR 34864). After the interim rule was published, 82 commenters addressed the duration of contract provision. Seventynine of these recommended change to the provision; 10 recommended a 1 year contract with an option for up to four yearly renewals (i.e. a maximum 5-year bidding cycle) and 69 recommended that the restriction on contract renewals be eliminated entirely so that each school food authority could determine the contract duration best suited to its own circumstances. However, as previously indicated, this issue will not be resolved in the final rewrite of Part 210 since the Department wishes to provide more specific notice to all members of the affected public of its intent to address the issue.

The concerns of commenters who recommended that the contract duration provision be lengthened or eliminated fall into two general categories. First, commenters indicated that the maximum 3-year duration of

management company contracts worked against the overall stability of school food service operations. They cited the adverse impact on food service employees of frequent potential or actual changes in management, the inability of companies to provide capital equipment to enhance food service operations in the absence of a suitable contract period over which to amortize such equipment, and the large amount of management company resources expended on developing proposals and responding to bid requests which could be better spent on Program operations. Secondly, commenters complained of the paperwork and administrative burden on school food authorities which results from the current 3-year maximum bidding cycle.

The Department is sensitive to commenter concerns on this issue, particularly those dealing with excessive administrative burden and paperwork. However, the Department is also committed to ensuring that the procurement of services from food service management companies is conducted in a manner that provides appropriate open and free competition consistent with the objectives and requirements of the NSLP. This commitment is in keeping with the procurement standards established for Federal grant programs under Office of Management and Budget Circular A-102. The Department believes there is sufficient merit in commenter concerns to warrant extending the maximum contract duration period from 3 to 5 years. The Department, therefore, is proposing to allow up to four annual renewals of food service management company contracts rather than the current two. This action would provide administrative relief to school food authorities and food service management companies alike and would contribute to a more stable food service environment in those school food authorities contracting with food service management companies. The Department believes that the maximum 5-year bidding cycle resulting from this proposal coupled with the current termination provision which allows a school food authority to cancel a food service management company contract for cause with 60-day notification, will continue to provide a climate conducive to free and open competition and satisfactory contract performance. The Department intends that this proposed provision would apply to contracts with food service management companies signed after the effective date of any final rule change.

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List of Subjects in 7 CFR Part 210

Food assistance programs, National School Lunch Program, Commodity School Program, Grant programs— Social programs, Nutrition, Children, Reporting and recordkeeping requirements, Surplus agricultural commodities.

Accordingly, Part 210 is amended as follows:

PART 210—NATIONAL SCHOOL LUNCH PROGRAM

1. The authority citation for Part 210 continues to read as follows:

Authority: Secs. 2–12, 60 Stat. 230, as amended; sec. 10, 80 Stat. 889, as amended; 84 Stat. 270; 42 U.S.C. 1751–1760, 1779, unless otherwise noted.

§ 210.16 [Amended]

2. In § 210.16, paragraph (d) is amended by removing the number "2" and adding, in its place, the number "4".

Date: August 19, 1987.

Anna Kondratas,

Administrator.

[FR Doc. 87-20053 Filed 8-31-87; 8:45 am] BILLING CODE 3410-30-M

Federal Crop Insurance Corporation

7 CFR Part 423

[Amdt. No. 1 (Doc. No. 4645S)]

Flaxseed Crop Insurance Regulations

AGENCY: Federal Crop Insurance Corporation, USDA.

ACTION: Proposed rule.

SUMMARY: The Federal Crop Insurance Corporation (FCIC) proposes to amend the Flaxseed Crop Insurance Regulations (7 CFR Part 423), effective for the 1988 crop year. The intended effect of this proposed rule is to maintain the effectiveness of the present Flaxseed Crop Insurance Regulations only through the 1987 crop year. It is proposed in a separate document that the provisions currently contained in this Part will be issued as an endorsement to the newly issued 7 CFR Part 401, General Crop Insurance Regulations as § 401.116, Flaxseed Endorsement, effective for the 1988 and succeeding crop years. 7 CFR Part 401 is a standard set of regulations and a master policy for insuring most crops which substantially reduces: (1) The time involved in amendment or revision; (2) the necessity of the present repetitious review process; and (3) the volume of paperwork processed by FCIC. The authority for the

promulgations of this rule is the Federal Crop Insurance Act, as amended.

DATE: Written comments, data, and opinions on this proposed rule must be submitted not later than October 1, 1987, to be sure of consideration.

ADDRESS: Written comments, data, and opinions on this proposed rule should be sent to Peter F. Cole, Office of the Manager, Federal Crop Insurance Corporation, Room 4090, South Building, U.S. Department of Agriculture, Washington, DC 20250. Written comments will be available for public inspection in the Office of the Manager, Room 4090, South Building, U.S. Department of Agriculture, Washington, DC during regular business hours, Monday through Friday.

FOR FURTHER INFORMATION CONTACT: Peter F. Cole, Secretary, Federal Crop Insurance Corporation, U.S. Department of Agriculture, Washington, DC 20250, telephone (202) 447–3325.

SUPPLEMENTARY INFORMATION: This action has been reviewed under USDA procedures established by Departmental Regulation 1512–1. This action does not constitute a review as to the need, currency, clarity, and effectiveness of these regulations under those procedures. The sunset review date established for these regulations is August 1, 1990.

E. Ray Fosse, Manager, FCIC, (1) has determined that this section is not a major rule as defined by Executive Order 12291 because it will not result in: (a) An annual effect on the economy of \$100 million or more; (b) major increases in costs or prices for consumers, individual industries, Federal, State, or local governments, or a geographical region; or (c) significant adverse effects on competition, employment, investment, productivity, innovation, or the ability of U.S.-based enterprises to compete with foreign-based enterprises in domestic or export markets; and (2) certifies that this action will not increase the Federal paperwork burden for individuals, small businesses, and other persons.

This action is exempt from the provisions of the Regulatory Flexibility Act; therefore, no Regulatory Flexibility Analysis was prepared.

This program is listed in the Catalog of Federal Domestic Assistance under No. 10.450.

This program is not subject to the provisions of Executive Order 12372 which requires intergovernmental consultation with State and local officials. See the Notice related to 7 CFR Part 3015, Subpart V, published at 48 FR 29115, June 24, 1983.

This action is not expected to have any significant impact on the quality of the human environment, health, and safety. Therefore, neither an Environmental Assessment nor an Environmental Impact Statement is needed.

Background

FCIC has published over 40 policies to cover insurance on that many different crops. Many of the regulations and policies contain identical language, which, if changed requires that over 40 different policies be changed, both in the Code of Federal Regulations (CFR) and the printed policy language. This repetition of effort is both inefficient and expensive. FCIC, therefore, has published in 7 CFR Part 401, one set of regulations and one master policy to contain that language which is identical in most of the policies and regulations.

As revisions on individual policies are necessary. FCIC proposes to publish a "crop endorsement" which will contain the language of the policy unique to that crop, and any exceptions to the master policy language necessary for that crop. When an endorsement is published as a section to Part 401, effective for a subsequent crop year, the present policy contained in a separate part of Chapter IV will be terminated at the end of the crop year then in effect.

In order to clearly establish that 7 CFR Part 423 will be effective only through the end of the 1987 crop year, FCIC herein proposes to amend the subpart heading of these regulations to specify that such will be the case.

It is proposed that the new Flaxseed Endorsement will be published as an endorsement to 7 CFR Part 401 (§ 401.116, Flaxseed Endorsement), and become effective for the 1988 and succeeding crop years. Upon final publication, the provisions of the Flaxseed Crop Insurance Regulations, now contained in 7 CFR Part 423, would be superseded. Therefore, FCIC proposes to amend the subpart heading to provide that 7 CFR Part 423 be effective for the 1986 and 1987 crop years only.

List of Subjects in 7 CFR Part 423

Crop insurance, Flaxseed.

Proposed Rule

Accordingly, pursuant to the authority contained in the Federal Crop Insurance Act, as amended (7 U.S.C. 1501 et seq.), the Federal Crop Insurance Corporation hereby passes to amend the Subpart heading to the Flaxseed Crop Insurance Regulations (7 CFR Part 423), as follows:

PART 423-[AMENDED]

1. The Authority citation for 7 CFR Part 423 continues to read as follows:

Authority: Secs. 506, 516, Pub. L. 75-430, 52 Stat. 73, 77, as amended (7 U.S.C. 1506, 1516).

The subpart heading in 7 CFR Part 423 is revised to read as follows:

Subpart—Regulations for the 1986 and 1987 Crop Years.

Done in Washington, DC, on July 24 1987. E. Ray Fosse,

Manager, Federal Crop Insurance Corporation.

[FR Doc. 87-20024 Filed 8-31-87; 8:45 am] BILLING CODE 3410-08-M

7 CFR Part 431

[Amdt. No. 1; (Doc. No. 4646S)]

Soybean Crop Insurance Regulations

AGENCY: Federal Crop Insurance Corporation, USDA. ACTION: Proposed rule.

SUMMARY: The Federal Crop Insurance Corporation (FCIC) proposes to amend the Sovbean Crop Insurance Regulations (7 CFR Part 431), effective for the 1988 crop year. The intended effect of this proposed rule is to maintain the effectiveness of the present Soybean Crop Insurance Regulations only through the 1987 crop year. It is proposed in a separate document that the provisions currently contained in this part will be issued as an endorsement to the newly issued 7 CFR Part 401, General Crop Insurance Regulations as § 401.117, Soybean Endorsement, effective for the 1988 and succeeding crop years. 7 CFR Part 401 is a standard set of regulations and a master policy for insuring most crops which substantially reduces: (1) The time involved in amendment or revision; (2) the necessity of the present repetitious review process; and (3) the volume of paperwork processed by FCIC. The authority for the promulgation of this rule is the Federal Crop Insurance Act, as amended.

DATE: Written comments, data, and opinions on this proposed rule must be submitted not later than October 1, 1987, to be sure of consideration.

ADDRESS: Written comments, data, and opinions on this proposed rule should be sent to Peter F. Cole, Office of the Manager, Federal Crop Insurance Corporation, Room 4090, South Building, U.S. Department of Agriculture, Washington, DC 20250. Written comments will be available for public.

inspection in the Office of the Manager, Room 4090, South Building, U.S. Department of Agriculture, Washington, DC during regular business hours, Monday through Friday.

FOR FURTHER INFORMATION CONTACT: Peter F. Cole, Secretary, Federal Crop Insurance Corporation, U.S. Department of Agriculture, Washington, DC, 20250, telephone (202) 447–3325.

SUPPLEMENTARY INFORMATION: This action has been reviewed under USDA procedures established by Departmental Regulation 1512–1. This action does not constitute a review as to the need, currency, clarity, and effectiveness of these regulations under those procedures. The sunset review data established for these regulations is October 1, 1990.

E. Ray Fosse, Manager, FCIC, (1) has determined that this action is not a major rule as defined by Executive Order 12291 because it will not result in: (a) An annual effect on the economy of \$100 million or more; (b) major increases in costs or prices for consumers, individual industries, Federal, State, or local governments, or a geographical region; or (c) significant adverse effects on competition, employment, investment, productivity, innovation, or the ability of U.S.-based enterprises to compete with foreign-based enterprises in domestic or export markets; and (2) certifies that this section will not increase the Federal paperwork burden for individuals, small businesses, and other persons.

This action is exempt from the provisions of the Regulatory Flexibility Act; therefore, no Regulatory Flexibility Analysis was prepared.

This program is listed in the Catalog of Federal Domestic Assistance under No. 10.450.

This program is not subject to the provisions of Executive Order 12372 which requires intergovernmental consultation with State and local officials. See the Notice related to 7 CFR Part 3015, Subpart V, published at 48 FR 29115, June 24, 1983.

This action is not expected to have any significant impact on the quality of the human environment, health, and safety. Therefore, neither an Environmental Assessment nor an Environmental Impact Statement is needed.

Background

FCIC has published over 40 policies to cover insurance on that many different crops. Many of the regulations and policies contain identical language, which, if changed requires that over 40 different policies be changed, both in the Code of Federal Regulations (CFR) and the printed policy language. This repetition of effort is both inefficient and expensive. FCIC, therefore, has published in 7 CFR Part 401, one set of regulations and one master policy to contain that language which is identical in most of the policies and regulations.

As revisions on individual policies are necessary. FCIC proposes to publish a "crop endorsement" which will contain the language of the policy unique to that crop, and any exceptions to the master policy language necessary for that crop. When an endorsement is published as a section to Part 401, effective for a subsequent crop year, the present policy contained in a separate part of Chapter IV will be terminated at the end of the crop year then in effect.

In order to clearly establish that 7 CFR Part 431 will be effective only through the end of the 1987 crop year. FCIC herein proposes to amend the subpart heading of these regulations to specify that such will be the case.

It is proposed that the new Soybean Endorsement will be published as an endorsement to 7 CFR Part 401 (§ 401.117, Soybean Endorsement), and become effective for the 1988 and succeeding crop years. Upon final publication, the provisions of the Soybean Crop Insurance Regulations, now contained in 7 CFR Part 431, would be superseded. Therefore, FCIC proposes to amend the subpart heading to provide that 7 CFR Part 431 be effective for the 1986 and 1987 crop years only.

List of Subjects in 7 CFR Part 431

Crop insurance, Soybean.

Proposed Rule

Accordingly, pursuant to the authority contained in the Federal Crop Insurance Act, as amended (7 U.S.C. 1501 et seq.), the Federal Crop Insurance Corporation hereby proposes to amend the subpart heading to the Soybean Crop Insurance Regulations (7 CFR Part 431), as follows:

PART 431-[AMENDED]

 The authority citation for 7 CFR Part 431 continues to read as follows:

Authority: Secs. 506, 516, Pub. L. 75–430, 52 Stat. 73, 77, as amended (7 U.S.C. 1506, 1516).

The subpart heading in 7 CFR Part
 431 is revised to read as follows:

Subpart—Regulations for the 1986 and 1987 Crop Years

Done in Washington, DC, on July 24, 1987. E. Ray Fosse,

Manager, Federal Crop Insurance Corporation.

[FR Doc. 87-20025 Filed 8-31-87;8:45 am] BILLING CODE 3410-08-M

Agricultural Marketing Service

7 CFR Parts 1136 and 1139

[Docket Nos. AO-309-A27 and AO-374-A11)]

Milk in the Great Basin and Lake Mead Marketing Areas; Extension of Time for Filing Exceptions on Proposed Amendments to Tentative Marketing Agreements and to Orders

AGENCY: Agricultural Marketing Service, USDA.

ACTION: Extension of time for filing exceptions to proposed rule.

SUMMARY: This notice extends the time for filing exceptions to a recommended decision issued July 14, 1987, concerning proposed amendments to the Great Basin and Lake Mead milk marketing orders. The Holstein-Friesian Association of America requested additional time to complete exceptions to the recommended decision.

DATE: Exceptions now are due on or before September 4, 1987.

ADDRESS: Exceptions (four copies) should be filed with the Hearing Clerk, Room 1079, South Building, United States Department of Agriculture, Washington, DC 20250.

FOR FURTHER INFORMATION CONTACT:

Constance M. Brenner, Marketing Specialist, USDA/AMS/Dairy Division, Order Formulation Branch, Room 2968, South Building, P.O. Box 96456, Washington, DC 20090-6456 [202] 447-(7183).

SUPPLEMENTARY INFORMATION: Prior documents in the proceeding:

Notice of Hearing: Issued February 6, 1986; published February 11, 1986 (51 FR 5070).

Proposed Suspension: Issued July 29, 1986; published August 4, 1986 (51 FR 27866).

Suspension Order: Issued September 2, 1986; published September 5, 1986 (51 FR 31759).

Recommended Decision: Issued July 14, 1987; published July 21, 1987 (52 FR 27372).

Notice is hereby given that the time for filing exceptions to the recommended decision with respect to the proposed amendments to the tentative marketing agreements and to the orders regulating the handling of milk in the Great Basin and Lake Mead marketing areas which was issued July 14, 1987, is hereby extended to September 4, 1987.

This notice is issued pursuant to the provisions of the Agricultural Marketing Agreement Act of 1937, as amended (7 U.S.C. 601 through 674), and the applicable rules of practice and procedure governing the formulation of marketing agreements and marketing orders (7 CFR Part 900).

List of Subjects in 7 CFR Parts 1136 and 1139

Milk marketing orders, Milk, Dairy products.

The authority citation for 7 CFR Parts 1136 and 1139 continues to read as follows:

Authority: Secs. 1-19, 48 Stat. 31, as amended; 7 U.S.C. 601-674.

Signed at Washington, DC, on August 26, 1987.

J. Patrick Boyle,

Administrator, Agricultural Marketing Service.

[FR Doc. 87-20020 Filed 8-31-87; 8:45 am]
BILLING CODE 3410-02-M

Farmers Home Administration

7 CFR Parts 1942, 1951, and 1955

Revision of Procedure to Service Community Program Loans Sold to the Private Sector With Servicing To Be Performed in the Private Sector

AGENCY: Farmers Home Administration, USDA.

ACTION: Proposed rule.

SUMMARY: The Farmers Home
Administration (FmHA) proposes to
amend its regulation to exclude the
servicing of loans sold without
insurance by FmHA to the private sector
with servicing to be performed in the
private sector. This action is necessary
to clearly establish servicing
responsibilities for such loans.

DATE: Comments must be received on or before September 16, 1987.

ADDRESSES: Submit written comments in duplicate to the Office of the Chief, Directives Management Branch, Farmers Home Administration, USDA, South Building, Room 6348, 14th and Independence Avenue SW., Washington, DC 20250. All written comments made pursuant to this notice will be available for public inspection on weekdays between the hours of 8:15 a.m. and 4:45 p.m. at the above address.

FOR FURTHER INFORMATION CONTACT: Bonnie S. Justice, Loan Officer, Community Facilities Division, Farmers Home Administration, U.S. Department of Agriculture, Room 6304, South Agriculture Building, Washington, DC 20250; telephone (202) 382–1490.

SUPPLEMENTARY INFORMATION: This proposed action has been reviewed under USDA procedures established in Departmental Regulation 1512-1, which implements Executive Order 12291, and has been determined to be "nonmajor" since the annual effect on the economy is less than \$100 million and there will be no significant increase in cost or prices for consumers; individual industries; Federal, State, or Local Government agencies; or geographic regions. Furthermore, there will be no adverse effects on competition, employment, investment, productivity, innovation, or on the ability of United States-based enterprises to compete with foreign-based enterprises in domestic or export markets.

This document has been reviewed in accordance with 7 CFR Part 1940,
Subpart G, "Environmental Program".
FmHA has determined that this action does not constitute a major Federal action significantly affecting the quality of the human environment and in accordance with the National Environmental Policy Act of 1969 Pub. L. 91–190, an Environmental Impact Statement is not required.

The Administrator, Farmers Home Administration, has determined that because of the limited scope of this action, and the requirements of the Omnibus Budget Reconciliation Act of 1986 (Pub. L. 99-509) (OBRA) and the Joint Resolution Making Continuing Appropriations for 1987 (Pub. L. 99–591), a fifteen-day comment period is necessary and adequate. The Farmers Home Administration is required to complete the sale of part of its Community Programs loan portfolio to the private sector no later than September 30, 1987. Loans will be sold to the private sector with servicing to be performed in the private sector as provided by OBRA. After the sale they will not be serviced by the Farmers Home Administration as Agency held loans. Pursuant to OBRA the Secretary of Agriculture will determine prior to sale that the private servicing arrangements will meet statutory requirements. The fifteen-day comment period will allow sufficient time for interested and affected persons to provide comments.

This change affects the following FmHA programs as listed in the Catalog of Federal Domestic Assistance and is subject to the provisions of Executive Order 12372 which requires

intergovernmental consultation with State and local officials. (7 CFR Part 3015, Subpart V, 38 FR 29112, June 24, 1983; 49 FR 22675, May 31, 1984; 50 FR 14088, April 10, 1985):

10.418 Water and Waste Disposal Systems for Rural Communities.10.423 Community Facilities Loans.

Discussion

FmHA is authorized by the Omnibus Budget Reconciliation Act to sell loans to the private sector with servicing to be performed in the private sector. This proposed action is to clearly establish servicing responsibilities for such loans and to clarify in FmHA's servicing regulations to show that those loans will be serviced in the private sector. This proposed action will provide that future changes to FmHA regulations will not be applicable to such loans.

List of Subjects

7 CFR Part 1942

Community development, Community facilities, Loan programs—Housing and community development, Loan security, Rural areas, Waste treatment and disposal—Domestic, Water supply—Domestic.

7 CFR Part 1951

Account servicing; Grant programs— Housing and community development; Reporting requirements; Rural areas; Subsidies.

7 CFR Part 1955

Foreclosure; Government acquired property; Government property management.

Accordingly, FmHA proposes to amend Chapter XVIII, Title 7, Code of Federal Regulations as follows:

PART 1942—ASSOCIATIONS

1. The authority citation for Part 1942 continues to read as follows:

Authority: 7 U.S.C. 1989; 16 U.S.C. 1005; 7 CFR 2.23; 7 CFR 2.70,

Subpart A—Community Facility Loans

2. § 1942.1 is amended by redesignating paragraph (c) as paragraph (d) and by adding a new paragraph (c) to read as follows:

§ 1942.1 General.

(c) Loans sold without insurance by FmHA to the private sector will be serviced in the private sector and will not be serviced under this subpart. The provisions of this subpart are not applicable to such loans. Future changes

to this subpart will not be made applicable to such loans.

PART 1951—SERVICING AND COLLECTIONS

3. The authority citation for Part 1951 continues to read as follows:

Authority: 7 U.S.C. 1989; 42 U.S.C. 1480; 5 U.S.C. 301; 7 CFR 2.23; 7 CFR 2.70.

Subpart E—Servicing of Community Program Loans and Grants

4. § 1951.201 is revised to read as follows:

§ 1951.201 Purpose.

This subpart prescribes the policies, authorizations, and procedures for servicing Community Water and Waste Disposal System loans and grants, Community Facility Loans, Industrial Development grants, loans for Grazing and other shift-in-land use projects, Association Recreation loans, Association Irrigation and Drainage loans, Watershed loans and advances, Resource Conservation and Development loans, Economic Opportunity Cooperative loans, loans to Indian Tribes and Tribal Corporations, loans to Timer Development Organizations, Rural Renewal loans and **Energy Impacted Area Development** Assistance Program grants. Loans sold without insurance by the Farmers Home Administration to the private sector will be serviced in the private sector and will not be serviced under this subpart. The provisions of this subpart are not applicable to such loans. Future changes to this subpart will not be made applicable to such loans.

Subpart O—Servicing Cases Where Unauthorized Loans(s) or Other Financial Assistance Was Received—Community and Insured Business Programs

5. § 1951.701 is revised to read as follows:

§ 1951.701 Purpose.

This subpart prescribes the policies and procedures for servicing Community and Business Program loans and/or grants made by Farmers Home Administration (FmHA) when it is determined that the borrower or grantee was not eligible for all or part of the financial assistance received in the form of a loan, grant, or subsidy granted, or any other direct financial assistance. It does not apply to guaranteed loans. Loans sold without insurance by the FmHA to the private sector will be serviced in the private sector and will

not be serviced under this subpart. The provisions of this subpart are not applicable to such loans. Future changes to this subpart will not be made applicable to such loans.

PART 1955-PROPERTY MANAGEMENT

6. The authority citation for Part 1955 continues to read as follows:

Authority: 7 U.S.C. 1989; 42 U.S.C. 1480; 5 U.S.C. 301; 7 CFR 2.23; 7 CFR 2.70.

Subpart A—Liquidation of Loans Secured by Real Estate and Acquisition of Real and Chattel Property

7. § 1955.1 is revised to read as follows:

§ 1955.1 Purpose.

This subpart delegates authority and prescribes procedures for the liquidation of Farmers Home Administration (FmHA) loans identified in § 1955.3 (d) and (e) of this subpart and acquisition of property by voluntary conveyance to the Government, by foreclosure of security instruments, by exercise of the Government's redemption rights, and certain other actions which result in acquisition of property by the Government. When FmHA elects to liquidate a guaranteed loan other than Business and Industrial (B&I) under the contract of guarantee, the liquidation will be completed according to this Subpart. Liquidations of guaranteed B&I loans will be effected upon direction from the Assistant Administrator, Community and Business Programs. For Community Programs and insured B&I actions involving loans secured by other than real or chattel property, the case will be forwarded to the National Office for prior review and guidance. Community Program loans sold without insurance by the FmHA to the private sector will be serviced in the private sector and will not be serviced under this subpart. The provisions of this subpart are not applicable to such loans. Future changes to this subpart will not be made applicable to such loans.

Subpart B-Management of Property

8. § 1955.51 is amended by adding a new paragraph (d) to read as follows:

§ 1955.51 Purpose.

(d) Community Program loans sold without insurance by the FmHA to the private sector will be serviced in the private sector and will not be serviced under this subpart. The provisions of this subpart are not applicable to such loans. Future changes to this subpart

will not be made applicable to such loans.

Date: August 26, 1987.

Vance L. Clark,

Administrator, Farmers Home Administration.

[FR Doc. 87-20079 Filed 8-31-87; 8:45 am]

7 CFR Parts 1951 and 1965

Security Servicing for Single Family Housing (SFH) Loans

AGENCY: Farmers Home Administration, USDA.

ACTION: Proposed rule.

SUMMARY: Farmers Home Administration (FmHA) proposes to amend its regulations regarding the sale of FmHA financed property by a borrower when the FmHA debt and authorized selling expenses exceeds the market value of the property. This action is being taken to expand the Agency's regulations on security servicing. The intended effect is to provide FmHA borrowers with the opportunity to voluntarily sell the property if necessary, while assisting the Agency in reducing the number of properties acquired into inventory through voluntary conveyance.

DATE: Comments must be submitted on or before November 2, 1987.

ADDRESSES: Submit written comments in duplicate to the Chief, Directives and Forms Management Branch, Farmers Home Administration, U.S. Department of Agriculture, Room 6348, South Agriculture Building, 14th Street and Independence Avenue, SW., Washington, DC 20250. All written comments made pursuant to this publication will be available for public inspection during regular work hours at the above address.

FOR FURTHER INFORMATION CONTACT:

David J. Villano, Senior Realty Specialist, Single Family Housing, Servicing and Property Management Division, FmHA, USDA, Room 5309, South Agriculture Building, Washington, DC 20250, Telephone (202) 382–1452.

SUPPLEMENTARY INFORMATION: This proposed rulemaking has been reviewed under USDA procedures established in Departmental Regulation 1512–1 which implements Executive Order 12291 and has been classified as "nonmajor." It will not result in an annual effect on the economy of \$100 million or more; a major increase in costs or prices for consumers, individual industries, Federal, State, or local government agencies, or geographic regions; or

significant adverse effects on competition, employment, investment, productivity, innovation, or on the ability of United States-based enterprises to compete with foreignbased enterprises in domestic or export markets

The SFH program is listed in the Catalog of Federal Domestic Assistance under No. 10.410—Low Income Housing Loans and No. 10.417—Very Low Income Housing Repair Loans and Grants. For the reasons set forth in the Final Rule related Notice(s) to 7 CFR Part 3015, Subpart V, this program is excluded from the scope of Executive Order 12372 which requires intergovernmental consultation with State and local officials.

This document has been reviewed in accordance with 7 CFR Part 1940, Subpart G. Environmental Program. It is the determination of FmHA that this action does not constitute a major Federal action significantly affecting the quality of the human environment, and, in accordance with the National Environmental Policy Act of 1969, Pub. L. 91–90, an Environmental Impact Statement is not required.

This proposed rule has been reviewed with regard to the requirements of the Regulatory Flexibility Act [5 U.S.C. 601 through 612]. The undersigned has determined and certified by signature of this document that this rule will not have a significant economic impact on a substantial number of small entities.

Current FmHA regulations [7 CFR Part 1965, Subpart C, § 1965.125(a)(2)] authorize FmHA officials to consent to the sale of SFH security property for less than the debt if the proposed selling price is at least equal to the current market value of the property as determined by FmHA. Where the purchaser is buying the property with personal funds or is obtaining credit from a source other than FmHA, and cash proceeds are available at closing. FmHA regulations further permit costs "* * * which the seller customarily or legally must pay to convey title and include but are not limited to: A real estate broker's commission, no more than three discount points to enable the buyer to obtain credit from another lender provided they are not being paid to reduce the purchaser's interest rate. real estate taxes, preparation of the deed, abstract fees, termite inspection, and deed or other revenue stamps." to be deducted from the (cash) sales proceeds before applying the remainder to the FmHA debt.

Said authority provides FmHA borrowers with an incentive to sell their house when there is no equity in the property. Due to the poor economy in many of the rural areas in which FmHA finances, property values have declined causing the FmHA debt to exceed market value. This authority has assisted many borrowers to sell their FmHA financed property where they have needed or desired to sell same due to employment, health, or other reasons.

A problem arises, however, when a FmHA borrower proposes to sell the property and an FmHA applicant desires to purchase the house and assume the existing FmHA debt. Where the FmHA debt exceeds market value, current FmHA regulations authorize the debt to be assumed at the market value. In these cases, there are no cash proceeds available at closing for the FmHA borrower/seller to pay necessary closing costs such as attorney fees, a real estate broker's commission, etc. This lack of cash proceeds generally precludes any such sales of an FmHA financed property to an FmHA program applicant, which is contrary to Agency policy.

If borrowers cannot sell their property, the next method to voluntarily liquidate their FmHA indebtedness is voluntary conveyance. In a voluntary conveyance, a borrower conveys title to the FmHA financed property to FmHA in exchange for a full or partial satisfaction of the debt. In most cases, FmHA will accept the voluntary conveyance and take title to the property. The agency is then responsible for making the property readily marketable through repairs, selling the property, and then paying all expenses incident to the sale such as transfer costs and a real estate broker's commission. This process is very lengthy and costly to the Government. Since FmHA eventually pays the selling expenses incident to the sale, we believe it would be proper, reasonable, rational, and in the Government's best interest to authorize payment of necessary selling expenses when the borrower proposes to sell the property and there are insufficient funds at the closing to pay same. Such a policy would be consistent with existing regulations which, as previously mentioned, authorize the Government to accept less than market value for payment of the indebtedness when cash

Due to the aforementioned reasons, FmHA has determined that payment by the Agency of authorized selling expenses when necessary to consummate a sale is reasonable and proper. Additionally, said action is necessary and rational as the Agency would be assisting its borrowers in

proceeds are available.

furthering the objectives of the program while reducing the number of properties acquired into inventory and the high costs associated with same.

Accordingly, FmHA proposes to expand and clarify its existing regulations to permit the Agency to pay for expenses on behalf of the borrower incident to the sale of their house when sufficient funds are not available to pay same. Further, consistent with existing Agency regulations regarding voluntary conveyances [See 7 CFR Part 1955, Subpart A, § 1955.10(c)(2)], settlement of a junior lien(s) is also authorized when determined to be in the Government's best financial interest.

List of Subjects

7 CFR Part 1951

Account servicing, Rent subsidies, Subsidies.

7 CFR Part 1965

Administrative practice and procedure, Loan programs, Housing and community development, Low and moderate income housing—Rental, Rural areas.

Therefore, as proposed, Chapter XVII of Title 7, Code of Federal Regulations, is amended as follows:

PART 1951—SERVICING AND COLLECTIONS

1. The authority citation for Part 1951 continues to read as follows:

Authority: 7 U.SC. 1989; 42 U.S.C. 1480; 5 U.S.C. 301; 7 CFR 2.23; 7 CFR 2.70.

Subpart M—Servicing Cases Where Unauthorized Loan or Other Financial Assistance Was Received—Single Family Housing

§ 1951.612 [Amended]

2. In § 1951.612(a)(1)(iii), the first sentence is amended by changing the reference "§ 1965.125(a)(4)" to "§ 1965.125(a)(3)."

PART 1965—REAL PROPERTY

3. The authority citation for Part 1965 continues to read as follows:

Authority: 7 U.S.C. 1989; 41 U.S.C. 2942; 42 U.S.C. 1480; 5 U.S.C. 301; 7 CFR 2.23; 7 CFR 2.70; 29 FR 14764; 33 FR 9850.

Subpart C—Security Servicing for Single Family Rural Housing Loans

4. In § 1965.125, paragraph (a)(3) is removed, paragraph (a)(4) is redesignated as (a)(3), and paragraph (a)(2) is revised to read as follows:

§ 1965.125 Liquidation

(a) * * *

(2) Consent to sale when the FmHA debt and authorized selling expenses exceed market value. If a borrower proposes to sell the property for an amount which will be insufficient to pay the FmHA debt, prior lien(s), if any, and authorized selling expenses, the County Supervisor will take a financial statement from the borrower on Form FmHA 431-3 "Household Financial Statement and Budget" to determine if they have reasonable ability to pay all or a substantial portion of the authorized selling expenses. In making this determination, consideration will be given to the borrower's moving and relocation expenses and the Government's prospects of acquisition of the property by voluntary conveyance or foreclosure and related costs of same. The County Supervisor will appraise the property and may consent to the sale if the proposed sale price is not less than the market value.

(i) Authorized selling expenses. Authorized selling expenses are those which the seller customarily or legally must pay to convey title and include but are not limited to: A real estate broker's commission which does not exceed the prevailing rate for the sale of similar property in the area, no more than three points to enable the buyer to obtain credit from another lender provided they are not being paid to reduce the purchaser's interest rate, real estate taxes, junior liens in the same manner as outlined in § 1955.10(c)(2) of Subpart A of Part 1955 of this chapter, preparation of the deed, abstract fees, termite inspection, and deed or other revenue stamps.

(ii) Closing the transaction. In no case will the borrower (seller) receive any cash proceeds from the sale.

Distribution of funds will be handled as follows:

(A) Where there are sufficient cash proceeds at closing, the entire sales proceeds, minus prior liens, if any, and authorized selling expenses, must be applied to the FmHA debt.

(B) Where cash proceeds are not available (such as in the case of an assumption) or are insufficient to pay authorized selling expenses, FmHA may pay said expenses necessary to consummate the transaction by preparation of Standard Form 1034, "Public Voucher For Purchases And Expenses Other Than Personal," and submission of Form FmHA 2024–1, "Miscellaneous Payment System," according to FmHA Instruction 2024–P (available in any FmHA office) and the respective Forms Manual Insert (FMI). Expenses so vouchered will be charged

to the borrower's (seller's) account as a recoverable cost item.

(iii) Release from liability. When consent under this paragraph is given, the County Supervisor is authorized to release the FmHA security instrument(s). When necessary to comply with State law, a State Supplement approved by OGC will prescribe procedures for releasing security instruments when the debt evidenced therein is not satisfied in full. Release of the borrower from liability for the deficiency is covered in § 1965.127(a)(4) and (b) of this subpart.

Dated: June 9, 1987. Vance L. Clark,

Administrator, Farmers Home Administration.

[FR Doc. 87–19584 Filed 8–31–87; 8:45 am] BILLING CODE 3410-07-M

DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Part 39

[Docket No. 87-CE-14-AD]

Petition of the Aircraft Owners and Pilots Association (AOPA) For Recision of Airworthiness Directive 87–08–08

AGENCY: Federal Aviation Administration (FAA), DOT, ACTION: Petition for rulemaking.

SUMMARY: This notice publishes for public comment a summary of the Aircraft Owners and Pilots Association (AOPA) petition dated may 28, 1987, as amended by their letter of August 3, 1987. This petition seeks the modification of Airworthiness Directive (AD) 87-08-08. The AD requires a wing spar inspection of certain Piper Aircraft Corporation Models PA-28 and PA-32 airplanes, and was issued as a result of a spar failure which occurred March 30, 1987. The petitioner contends that the service histories of the identified airplanes having spar cracks played an integral part in the development of the fatigue and the FAA should consider requiring compliance only on airplanes with a history of extensive low-level flight or other abnormal flight conditions. The petitioner further requests that the AD compliance times be modified or adjusted to reduce scheduling conflicts since no cracks have been reported on airplanes with less than 6100 hours.

Publication of this notice is not intended to affect the legal status of the petition or its final disposition.

DATE: Comments must be received on or before October 1, 1987.

FOR FURTHER INFORMATION CONTACT:

Raymond Boice, Aerospace Engineer, Aircraft Certification Division, 601 East 12th Street, 1656 Federal Building, Kansas City, Missouri 64106, Telephone (816) 374–5934.

ADDRESSES: Sent comments on this petition in triplicate to: Federal Aviation Administration, Office of the Regional Counsel, Attn: Rules Docket, Docket No. 87-CE-14-AD, 601 East 12th Street, 1558 Federal Building, Kansas City, Missouri 64106. Comments may be inspected in Room 1558 weekdays, except Federal holidays, between the hours of 7:30 a.m. and 4:00 p.m. In addition, the FAA is maintaining an information docket of comments in the Office of the Chief Counsel, Federal Aviation Administration, 800 Independence Avenue, SW., Washington, DC 20591. Comments may be inspected in Room 915G weekdays, except Federal holidays, between the hours of 8:30 a.m. and 5:00 p.m.

SUPPLEMENTARY INFORMATION:

Comments Invited

Interested persons are invited to submit such written data, views, or arguments on the petition as they may desire. Communications should identify the docket and petition number and be submitted in triplicate to the Office of Regional Counsel, Kansas City, Missouri, at the above address. All communications received on or before the closing date for comments will be considered before taking action on the petition. All comments will be available for examination in the FAA docket.

Interested persons may obtain a copy of the petition by contacting the person listed above in the paragraph entitled "FOR FURTHER INFORMATION CONTACT.."

Although this notice refers to the contents of the petition as received by the FAA, it should be understood that the purpose of the reference is to receive public comments in accordance with FAA procedures governing petitions for rulemaking, and it does not propose a regulatory rule for adoption or recision, represent an FAA position, or otherwise commit the FAA on the merits of the petition. The FAA intends to consider the merits of the proposal after it has had an opportunity to evaluate the petition matters presented and all comments received from the public.

The Petition

Accordingly, the Federal Aviation Administration publishes this notice for public comment on the AOPA petition for recision of AD 87–08–08. Issued in Kansas City, Missouri on August 14, 1987.

Paul K. Bohr,

Director, Central Region. [FR Doc. 87–19995 Filed 8–31–87; 8:45 am] BILLING CODE 4910–13–M

FEDERAL COMMUNICATIONS COMMISSION

47 CFR Parts 36 and 67

[CC Docket Nos. 78-72, 80-286 and 86-297; FCC 87-272]

Common Carrier Services; MTS and WATS Market Structure; Amendment of the Commission's Rules and Establishment of a Joint Board

AGENCY: Federal Communications Commission.

ACTION: Supplemental notice of proposed rulemaking.

SUMMARY: The Commission has issued a Supplemental Notice of Proposed Rulemaking seeking comment and data on the appropriate allocation method and recovery mechanism for marketing expenses (See 45 FR 41459, June 19. 1980). In a Memorandum Opinion and Order on Reconsideration, summarized elsewhere in this volume, the Commission granted, in part petitions for reconsideration of its decision to exclude access revenues from the allocation factor for marketing expenses. The Commission adopted a new § 36.372 of its rules, which will allocate marketing expenses on the basis of current billings, effective January 1, 1988.

DATES: Comments and data must be filed on or before September 28, 1987, and reply comments on or before October 13, 1987.

ADDRESS: Federal Communications Commission, Washington, DC 20554.

FOR FURTHER INFORMATION CONTACT: Cindy Schonhaut, Special Counsel Federal-State Joint Board Matters, Accounting and Audits Division, Common Carrier Bureau, at (202) 632–7500.

SUPPLEMENTARY INFORMATION: This is a summary of the Commission's Supplemental Notice of Proposed Rulemaking, CC Docket Nos. 78–72, 80–286 and 86–297, FCC 87–272; adopted August 14, 1987, and released August 18, 1987.

The full text of Commission decisions are available for inspection and copying during normal business hours in the FCC Dockets Branch (Room 230), 1919 M Street NW., Washington, DC. The

complete text of this decision may also be purchased from the Commission's copy contractors, International Transcription Service, 2100 M Street NW., Suite 140, Washington, DC 20037, (202) 857–3800.

Summary of Supplemental Notice of Proposed Rulemaking

1. The Joint Board in CC Docket No. 86–297 recommended that the Commission adopt a new separations manual intended to conform separations procedures to the recently revised Uniform System of Accounts (USOA) and to simplify the separations process. The Commission adopted the new Separations Manual in April 1987 which will be codified as the new Part 36 of its rules and which will become effective

January 1, 1988.2

2. Under the current separations procedures of Part 67 of the Commission's rules, the advertising and sales expenses in the current USOA Accounts 642 and 643 are allocated between the jurisdictions on the basis of current billing for local and toll services (excluding certain billings for nonaffiliated companies and those in connection with intercompany settlements).3 Those expenses will be included in the new Account 6610 in the revised USOA. Under new separations procedures recommended by the Joint Board and adopted by the Commission, billings for access charges will be excluded from the allocation factor for Account 6610 marketing expenses.

3. The Federal Communications Commission granted, in part, reconsideration of its decision to adopt revisions of the Separations Manual, new Part 36 of its rules, recently recommended by the Federal-State Joint Board. The Commission decided to reconsider its decision to exclude access revenues from the allocation factor for marketing expenses. Thus, as an interim measure, in the new § 36.372 of the Commission's rules (effective January 1, 1988) marketing expenses will be allocated on the basis of current billings. The Commission did not act, at this time on other reconsideration issues.

4. The Commission referred a permanent resolution of this issue to the Joint Board in CC Docket No. 80–286.

The Commission specified certain question for comment and requested data regarding the allocation of marketing expenses. The Commission asked the Joint Board to recommend a permanent resolution of this issue by April 1, 1988. The Commission also proposed to revise the Part 69 access apportionment rules for Account 6610 expenses to conform to any separations rule that may be adopted and invited the Joint Board in CC Docket No. 80–286 to recommend an appropriate rule.

5. The Commission requested that LECs submit data, as specified in Appendix B of the Notice, to aid the Joint Board and the Commission in the evaluation of alternative allocation methods. It also requested that LECs specifically identify their Account 6610 marketing activities that are related to a specific jurisdiction. The Commission requested that parties comment on the need for changes in the procedures for the allocation and recovery of expenses in Account 6610, including the desirability of excluding all or some portion of access revenues from the allocation factor or factors (e.g., subscriber line or carrier common line charge revenues,) and the ability of carriers to directly assign some portion of Account 6610 expenses to the state or interstate jurisdictions. It also sought specific proposals for alternative allocation factors and cost recovery mechanisms that, for example, may not be revenue-based, or may incorporate a weighting factor or phase-in approach.

Comments

6. Interested parties may file comments and data on the issues discussed above on or before September 28, 1987, and reply comments on or before October 13, 1987.4

Regulatory Flexibility Act

7. We certify that the Regulatory Flexibility Act 5 is not applicable to the rule changes we are proposing in this proceeding. In accordance with the provisions of section 605 of the Act, a copy of this certification will be sent to the Chief Counsel for Advocacy of the Small Business Administration at the time of publication of a summary of this Order and Notice in the Federal Register. As part of our analysis of the proposals received in response to the Notice, however, the Commission will consider the impact of proposals on small telephone companies, i.e., those serving 50,000 or fewer lines.6

Paperwork Reduction Act

8. We have analyzed the rules adopted and proposed herein with respect to the Paperwork Reduction Act of 1980 7 and have concluded (tentatively with respect to the proposals) that they will not impose new or modified information collection requirements on the public. In addition, the Notice presented herein is a general solicitation of comments from the public and, as such, does not constitute a collection of information.8 All comments will be considered in this proceeding. Parties need not specifically respond to the data request for their comments to be considered. Therefore, implementation of the requirements adopted and proposed herein will not be subject to approval by the Office of Management and Budget as prescribed by the Act.

Ex Parte Contacts

9. For purposes of the non-restricted notice and comment rulemaking proceeding which we are hereby initiating, members of the public are advised that ex parte contacts are permitted from the time the Commission adopts a notice of proposed rulemaking until the time a public notice is issued stating that a substantive disposition of the matter is to be considered at a forthcoming meeting or until a final order disposing of the matter is adopted by the Commission, whichever is earlier. In general, an ex parte presentation is any written or oral communication (other than formal written comments, pleadings and oral arguments) between a person outside the Commission and a Commissioner, Joint Board member, or a member of the staff of the Commission or the Joint Board, that addresses the merits or outcome of the proceeding. Any person who submits a written ex parte presentation must serve a copy of that presentation on the Commission's Secretary for inclusion in the public record. Any person who makes an oral ex parte presentation presenting data or

¹ Amendment of Part 67 (New Part 36) of the Commission's Rules and Establishment of Federal-State Joint Board, CC Docket No. 86–297, FCC 87]–4, released April 8, 1987, 2 FCC Rod 2,582 (1987) (Recommended Decision and Order).

^{*} MTS and WATS Market Structure, Amendment of Part 67 (New Part 36) of the Commission's Rules and Establishment of a Federal-State Joint Board, CC Docket Nos. 78-72, 80-286 and 86-297, FCC 87-134, released May 1, 1987, 2 FCC Rcd 2,639 (1987) (Report and Order).

^{3 47} CFR 67.363.

^{*} See 47 CFR 1.415 and 1.419.

⁵ U.S.C. 603.

⁶ Because of the nature of local exchange and access service, this Commission concluded that

small telephone companies are dominant in their fields of operation and, therefore, are not small entities as defined by the Regulatory Flexibility Act. See MTS and WATS Market Structure, 93 FCC 2d 241, 338–39 (1983). Thus, this Commission is not required by the terms of the Act to apply the formal procedures set forth therein. This Commission and the Joint Board are nevertheless committed to reducing the regulatory burdens on small telephone companies whenever possible consistent with our other public interest responsibilities. Accordingly, we have chosen to utilize, on an Informal basis, appropriate Regulatory Flexibility Act procedures to analyze the effect of the proposed regulations on small telephone companies.

^{7 44} U.S.C. 501.

^{*} See 5 CFR 1320.7(k)(4).

arguments not already reflected in that person's written comments, memoranda, or other previous filings in this proceeding, shall provide, on the day of the oral presentation, a written memorandum to the Secretary of the Commission (with a copy to the Commissioner, Joint Board member, or staff member involved), which summarizes the data and arguments in the ex parte presentation. Each such written memorandum must state on its face that the Secretary has been served, and must also state by docket number the proceeding to which it relates.

10. In reaching their decisions, this Commission and the Joint Board may consider information and ideas not contained in the comments, provided that such information or a writing indicating the nature and source of such information is placed in the public record and providing that the reliance of this Commission or the Joint Board on such information is noted in the Recommended Decision and Order or the Report and Order.

11. For Joint Board actions, special exparte rules apply. 10 For Joint Board actions, all written materials which are not filed in accordance with a pleading cycle established by the Joint Board shall be accompanied by a Petition for

shall be accompanied by a Petition for Leave to File showing cause why the material should be considered by the Joint Board. The Joint Board will not consider any filing made outside the authorized pleading cycle and received by the Commission less than fifteen days 11 in advance of a Joint Board meeting at which the Joint Board is to consider the subject matter of that filing. Written ex parte presentations, as defined by the Commission's rules, need not be accompanied by a Petition for Leave to File and may be received in the discretion of the Joint Board member or staff personnel involved. No written ex parte presentations, however, shall be made during the fifteen day period immediately preceding a Joint Board meeting except in response to an inquiry initiated by a member of the Joint Board or its staff.

Ordering Clause

 Accordingly, it is ordered, That the Joint Board in CC Docket No. 80–286

⁹ See generally Amendment of Subpart H, Part 1 of the Commission's Rules and Regulations Concerning Ex Parte Communications and shall review the comments, proposals and data and prepare recommendations to this Commission on the issues raised herein.

List of Subjects in 47 CFR Parts 36 and 67

Communications common carrier, Telephone, Uniform system of accounts, Reporting and recordkeeping requirements, Jurisdictional separations procedures.

This action is taken pursuant to 47 U.S.C. 154 (i) and (j), 201, 202, 205, 218, 221(c), 403 and 410.

Federal Communications Commission. William J. Tricarico,

Secretary.

[FR Doc. 87-19260 Filed 8-31-87; 8:45 am]

DEPARTMENT OF THE INTERIOR

Fish and Wildlife Service

50 CFR Part 17

Endangered and Threatened Wildlife and Plants; Proposed Endangered Status for the James Spinymussel

AGENCY: Fish and Wildlife Service, Interior.

ACTION: Proposed rule.

SUMMARY: The Service proposes to determine endangered status for the James spinymussel (Pleurobema collina). Critical habitat is not proposed. This species survives only in a few headwater streams of the James River in Virginia and West Virginia. This action is being taken because: (1) The range and numbers of this freshwater mussel have been drastically reduced to about 5-10% of historic levels, and (2) the few drainages that continue to support the species are subject to threats including invasion of essential habitats by the exotic Asiatic clam (Corbicula fluminea) and potential water quality degradation by agricultural and silvicultural runoff, effluent from sewage treatment plants, and chemical spills. This proposal, if made final, would implement Federal protection provided by the Endangered Species Act of 1973, as amended. The Service is requesting data and comments from the public on this proposal.

DATES: Comments from all interested parties must be received by November 2, 1987. Public hearing requests must be received by October 16, 1987.

ADDRESSES: Comments and materials concerning this proposal should be sent to the Annapolis Field Office, U.S. Fish and Wildlife Service, 1825B Virginia Street, Annapolis, Maryland 21401. Comments and materials received will be available for public inspection, by appointment, during normal business hours at the above address.

FOR FURTHER INFORMATION CONTACT: Mr. G. Andrew Moser at the above address (301/269-6324).

SUPPLEMENTARY INFORMATION:

Background

The James spinymussel was first discovered in the Calfpasture River, Rockbridge County, Virginia, by T. A. Conrad in 1836 (Conrad 1846). The species was orginally described by Conrad (1837) as *Unio collinus*. It has been subsequently placed in different genera by various workers. Names that refer to this species are listed in the following abbreviated synonymy:

Unio collinus Conrad, 1936: Plate 36, Figure 2.

Margaron (Unio) collinus (Conrad).— Lea 1852:23.

Alasmidonta collina (Conrad).— Simpson 1900:669.

Canthyria collina (Conrad).—Prierson 1927:1946; Stansberry 1971:14; Clarke and Neves 1984; Zeto and Schmidt 1984:147.

Elliptio (Canthyria) collina (Conrad).— Morrison 1955:20.

Pleurobema collina (Conrad).—Boss and Clench 1967:45; Heard 1970:27; Burch 1975:12.

Pleurobema (Lexingtonia) collina (Conrad).—Johnson 1970:300.

Fusconaia (Lexingtonia) collina (Conrad).—Johnson and Clarke 1983:296.

The Service recognized the James spinymussel under the name Fusconaia collina in the Review of Invertebrate Wildlife for Listing as Endangered or Threatened Species (49 FR 21675; May 22, 1984). Clarke and Neves (1984) subsequently determined that the James spinymussel uses only its outer gills to brood glochidia and is therefore not a Fusconaia, which are currently understood to use all four gills to brood glochidia. Clarke and Neves (1984) suggested placement of the species in the genus Canthyria, because of the presence of spines on the shell and some characters of the soft anatomy. The Service believes that until further review and evaluation clarifies the taxonomic significance of these characters, the James spinymussel should be recognized under the more established name Pleurobema collina.

The Service's Review of Invertebrate Wildlife included this species under the common name "Virginia spiny mussel."

Persentations in Commission Proceedings, GEN. Docket No. 86–225, 2 PCC Rcd 3,011 (1987). ¹⁰Amendment of Part 67 of the Commission's Rules and Establishment of a Joint Board, CC Docket No. 80–286, 89 PCC 2d 36 (1982).

¹¹ In calculating this fifteen day period, neither the day on which the material is filed nor the day on which the Joint Board meeting is scheduled shall be counted.

The Service is following the list of common names by Turgeon et al. (in press) in now using the name James

spinymussel.

The shells of juvenile James spinymussels usually bear one to three short but prominent spines on each valve. The shells of adults usually lack spines. The foot and mantle of the adult are conspicuously orange and the mantle is darkly pigmented in a narrow band around and within the edges of the branchial and anal openings.

Aside from the James spinymussel, only two other freshwater spined mussels are known to exist: Elliptio (Canthyria) spinosa, a large-shelled and long-spined species know only from the Altamaha River system in Georgia, and Eliptio (Canthyria) steinstansana, a species with intermediate shell size and spine length found only in the Tar River in North Carolina. The latter species was listed as endangered on June 27, 1985 (50 FR 26575). The James spinymussel has a smaller shell and shorter spines than these other two

species.

The James spinymussel has been collected on sand and mixed sand and gravel substrates, generally in areas of slow to moderate current and relatively hard water. Like other freshwater mussels, it feeds by filtering food particles from the water, a characteristic that makes it particularly susceptible to detrimental effects of water-borne pollutants. P. collina also shares with other freshwater mussels a complex reproductive cycle in which the mussel larvae attach for a short time to a fish host. The life span, time of spawning, host fish species, and many other aspects of the life history of P. collina are still unknown.

Collection records indicate that the James spinymussel was once widely distributed in the James River drainage upstream of Richmond. All pre-1983 records for the species are from Virginia (Clarke and Neves 1984). They include: the James River, main stem, in Rockbridge, Botetourt, Fluvanna, Buckingham, Goochland and Cumberland Counties; the Rivanna River in Fluvanna County; Mill Creek in Bath County; the Calfpasture River in Rockbridge County, and Johns Creek in Craig County. The James spinymussel was first reported from West Virginia in 1984 (Zeto and Schmidt 1984). According to a recent Service-funded survey of the James River drainage, this mussel is now restricted to Craig and Johns Creeks in Craig and Botetourt Counties, Virginia, and Potts Creek in Monroe County, West Virginia (Clarke and Neves 1984).

Although it is probable that the decline of the James spinymussel began with municipal growth and industrialization of cities and towns in the James River watershed, much of the decline has occurred in the last 20 years. The species remained in much of its historic range through the mid-1960's, but has since disappeared from the majority of known sites. It now appears to be extirpated from 90-95% of its historic range, with survival documented only in three headwater creeks in the James River drainage. This restricted distribution makes the species vulnerable to threats including water quality perturbations, disease, and displacement by expanding populations of the exotic Asiatic clam (Corbicula fluminea).

In the Federal Register on May 22, 1984 (49 FR 21675), the James spinymussel was included in category 2 of the Service's Review of Invertebrate Wildlife. Category 2 comprises those taxa for which proposed listing is possibly appropriate but for which conclusive data on biological vulnerability are not available to support a proposed rule. Additional data, including a Service-funded status survey (Clarke and Neves 1984) have provided the data needed to support a

listing proposal.

Summary of Factors Affecting the Species

Section 4(a)(1) of the Endangered Species Act (16 U.S.C. 1531 et seq.) and regulations (50 CFR Part 424) promulgated to implement the listing provisions of the Act set forth the procedures for adding species to the Federal Lists. A species may be determined to be an endangered or threatened species due to one or more of the five factors described in section 4(a)(1). These factors and their application to the James spinymussel (Pleurobema collina) are as follows:

A. The Present or Threatened Destruction, Modification, or Curtailment of its Habitat or Range

Results of a recent Service-funded survey of the James River drainage (Clarke and Neves 1984) indicate that the James spinymussel exists only in Craig and Johns Creeks in Craig and Botetourt Counties, Virginia, and a short reach of Potts Creek in Monroe County, West Virginia. This represents a very significant reduction (90-95%) in known range, as historic records indicate that the species was once found throughout much of the James River drainage upstream of Richmond.

Habitat modification has been a major factor in the James spinymussel's abrupt

decline. Adverse habitat changes including dam construction, industrial pollution, chemical spills, channelization, and sewage discharges have occurred at various locations within the species' historic range in the James River drainage. Current threats to habitat in the Craig/Johns Creek and Potts Creek watersheds include the following:

(1) Effluent discharges and accidental discharges of chlorine or raw sewage from the sewage treatment plant at New

Castle, Virginia;

(2) Erosion and siltation resulting from logging operations in the upper Craig Creek Watershed;

(3) Toxic chemical spills;

- (4) Agricultural runoff including pesticides and fertilizers;
 - (5) Channelization.

B. Overutilization of Commercial, Recreational, Scientific, or Educational Purposes

Although collection was probably an insignificant factor in this species' decline, it is becoming a problem now that the species is rare. Because additional interest in the spinymussel is expected to be generated by the listing process, the Service is concerned that this problem may worsen in the future.

C. Disease or Predation

There is no evidence that disease or predation has been a problem for the James spinymussel. However, extensive mussel dieoffs, possibly caused by a yet unknown disease, have occurred recently in the rivers of southwest Virginia, in the Tar River in North Carolina, and in numerous other locations. The Tar River dieoff, discovered in May 1986, was particularly severe, killing an estimated 75% of all mussels in the affected beds (R. Neves personal communication). Should such an outbreak occur in the Craig Creek or Potts Creek drainages, it would pose a very serious threat to the James spinymussel because of the species' restricted range.

D. The Inadequacy of Existing Regulatory Mechanisms

Virginia State law (section 29-113) requires a permit for the scientific collection of freshwater mussels. However, this State law is difficult to enforce and does not protect the species' habitat from the potential impacts of Federal projects. Federal listing would provide protection for the species under the Endangered Species Act by requiring a Federal permit to take the species and requiring Federal agencies to consult with the Service when projects they

fund, authorize, or carry out may affect the species.

E. Other Natural or Manmade Factors Affecting its Continued Existence

Much of the James River drainage has become infested by the Asiatic clam (Corbicula fluminea), a species introduced accidentally from Asia. Competition from this non-native species may be a principal cause of the James spinymussel's decline. Population densities of C. fluminea in excess of 1000 individuals per square meter (about 93 per square foot) have been reported in the James River downstream of Richmond (Diaz 1974). Because of the Asiatic clam's high population densities, its feeding activity may significantly reduce the availability of phytoplankton needed by the spinymussel for food and may interfere with reproduction of the spinymussel by filtering its sperm from the water column (Clarke 1981). Clarke and Neves (1984) consider the temporal correlation between the disappearance of downstream populations of the James spinymussel and the appearance and proliferation of the Asiatic clam to be clear evidence that the spread of Corbicula is one of the chief causes of the spinymussel's decline.

The Service has carefully assessed the best scientific and commercial information available regarding the past, present, and future threats faced by this species in determining to propose this rule. Based on this evaluation, the preferred action is to list the James spinymussel as endangered. The mussel's small population and restricted distribution make it vulnerable to pollution events, disease, and competition from exotic species; its range has greatly narrowed within the immediate past; therefore, threatened status would not be appropriate. The reasons for not proposing critical habitat for this species are discussed below in the "Critical Habitat" section.

Critical Habitat

Section 4(a)(3) of the Act, as amended. requires that to the maximum extent prudent and determinable, the Secretary designate critical habitat at the time a species is determined to be endangered or threatened. The Service finds that designation of critical habitat is not prudent for the James spinymussel at this time. This rare mussel is very unusual, being one of only three known species of spined freshwater mussels. There is a small but significant demand by collectors for this species. Because of this, the Service believes a detailed description of the species' habitat, required as part of any critical habitat designation, could increase the species'

vulnerability to illegal taking and increase law enforcement problems. Therefore, it would not be prudent to designate critical habitat for this species. Doing so would draw attention to the habitats supporting the James spinymussel and risk depletion of an already limited population.

Available Conservation Measures

Conservation measures provided to species listed as endangered or threatened under the Endangered Species Act include recognition. recovery actions, requirements for Federal protection, and prohibitions against certain practices. Recognition through listing encourages and results in conservation actions by Federal, State, and local governments and private agencies, groups, and individuals. The Endangered Species Act provides for possible land acquisition and cooperation with the States and requires that recovery actions be carried out for all listed species. Such actions are initiated by the Service following listing. The protection required of Federal agencies and the prohibitions against taking and harm are discussed, in part,

Section 7(a) of the Act, as amended, requires Federal agencies to evaluate their actions with respect to any species that is proposed or listed as endangered or threatened and with respect to its critical habitat, if any is being designated. Regulations implementing this interagency cooperation provision of the Act are codified at 50 CFR Part 402. Section 7(a)(4) requires Federal agencies to confer informally with the Service on any action that is likely to jeopardize the continued existence of a proposed species or result in destruction or adverse modification of proposed critical habitat. If a species is listed subsequently, section 7(a)(2) requires Federal agencies to ensure that activities they authorize, fund, or carry out are not likely to jeopardize the continued existence of such a species or to destroy or adversely modify its critical habitat. If a Federal action may affect a listed species or its critical habitat, the responsible Federal agency must enter into formal consultation with the Service.

Federal activities that could impact the James spinymussel and its habitat in the future include, but are not limited to, the following: issuance of permits for mineral exploration, timber sales, recreational development, stream alterations, road and bridge construction and maintenance, and implementation of forest management plans. It has been the experience of the Service that the large majority of section 7 consultations

are resolved so that the species is protected and the project can continue.

The Act and its implementing regulations found at 50 CFR 17.21 set forth a series of general prohibitions and exceptions that apply to all endangered wildlife. These prohibitions, in part, make it illegal for any person subject to the jurisdiction of the United States to take, import or export, ship in interstate commerce in the course of a commercial activity, or sell or offer for sale in interstate or foreign commerce any listed wildlife species. It also is illegal to possess, sell, delivery, carry, transport, or ship any such wildlife that has been illegally taken. Certain exceptions apply to agents of the Service and State conservation agencies.

Permits may be issued to carry out otherwise prohibited activities involving endangered wildlife species under certain circumstances. Applicable regulations governing permits are at 50 CFR 17.22 and 17.23. Such permits are available for scientific purposes, to enhance the propagation or survival of the species, and/or for incidental take in connection with otherwise lawful activities.

Public Comments Solicited

The Service intends that any final action resulting from this proposal will be accurate and as effective as possible. Therefore, any comments or suggestions from the public, other concerned governmental agencies, the scientific community, industry, or any other interested party concerning any aspect of these proposed rules are hereby solicited. Comments particularly are sought concerning:

 Biological, commercial trade, or other relevant data concerning any threat (or lack thereof) to the James spinymussel;

(2) The location of any additional populations of the James spinymussel and the reasons why any habitat should or should not be determined to be critical habitat as provided by section 4 of the Act;

(3) Additional information concerning the range and distribution of this species; and

(4) Current or planned activities in the subject area and their possible impacts on the James spinymussel.

Final promulgation of the regulations on the James spinymussel will take into consideration the comments and any additional information received by the Service, and such communications may lead to adoption of final regulations that differ from this proposal.

The Endangered Species Act provides for a public hearing on this proposal, if requested. Requests must be filed within 45 days of the date of the proposal. Such requests must be made in writing (see ADDRESSES section).

National Environmental Policy Act

The Fish and Wildlife Service has determined that an Environmental Assessment, as defined by the National Environmental Policy Act of 1969, need not be prepared in connection with regulations adopted pursuant to section 4(a) of the Endangered Species Act of 1973, as amended. A notice outlining the Service's reasons for this determination was published in the Federal Register on October 25, 1983 (48 FR 49244).

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Author

The primary author of this proposed rule is G. Andrew Moser, Annapolis Field Office, U.S. Fish and Wildlife Service, 1825B Virginia Street, Annapolis, Maryland 21401 (301/269-

List of Subjects in 50 CFR Part 17

Endangered and threatened wildlife, Fish, Marine mammals, Plants (agriculture).

Proposed Regulation Promulgation

Accordingly, it is hereby proposed to amend Part 17, Subchapter B of Chapter I, Title 50 of the Code of Federal Regulations, as set forth below:

PART 17-[AMENDED]

1. The authority citation for Part 17 continues to read as follows:

Authority: Pub. L. 93-205, 87 Stat. 884; Pub. L. 94-359, 90 Stat. 911; Pub. L. 95-632, 92 Stat. 3751; Pub. L. 96-159, 93 Stat. 1225; Pub. L. 97-304, 96 Stat. 1411 (16 U.S.C. 1531 et seq.).

2. It is proposed to amend § 17.11(h) by adding the following, in alphabetical order under Clams, to the List of Endangered and Threatened Wildlife:

§ 17.11 Endangered and threatened wildlife.

(h) * * *

Species			Vertebrate	3 12 1 ET	The same	Critical	Special
Common name	Scientific name	Historic range	population where endangered or threatened	Status	When listed	Critical habitat	Special rules
CLAMS	The second second			-		180	
Spinymussel, James (= Virginia spiny mussel),	Pleurobema (= Fusconaia) collina	U.S.A. (VA,WV)	NA	E		NA	NA

Dated: August 3, 1987.

Susan Recce.

Acting Assistant Secretary for Fish and Wildlife and Parks.

[FR Doc. 87-20022 Filed 8-31-87; 8:45 am]

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DEPARTMENT OF COMMERCE

National Oceanic and Atmospheric Administration

50 CFR Parts 611 and 675

[Docket No. 70878-7178]

Foreign Fishing; Groundfish of the Bering Sea and Aleutian Islands Area

AGENCY: National Marine Fisheries Service (NMFS), NOAA, Commerce.

ACTION: Proposed rule.

SUMMARY: NOAA issues a proposed rule

to implement Amendment 11 to the Fishery Management Plan for the Groundfish Fishery in the Bering Sea and Aleutian Islands Area (FMP) which is pending approval by the Secretary of Commerce (Secretary). The amendment would (1) establish a split season apportionment of pollock for U.S. vessels working in joint ventures with foreign processing vessels (JVP), and (2) change the definition of prohibited species. These measures are intended to respond to biological, economic and administrative problems identified by the North Pacific Fishery Management Council (Council)

In addition, NOAA is proposing to change the defintion of directed fishing.

The proposed regulations to implement Amendment 11 and the additional proposed regulatory change are necessary for conservation and management of the groundfish resources in the Bering Sea and Aleutian Islands (BSAI) area and for the orderly conduct of the groundfish fisheries

DATE: Comments on the amendment, proposed rule and supporting documents, especially the environmental assessment and regulatory impact review/initial regulatory flexibility analysis (EA/RIR/IRFA), are invited until October 15, 1987.

ADDRESS: Comments should be addressed to Robert W. McVey, Director, Alaska Region (Regional Director), National Marine Fisheries Service, P.O. Box 21668, Juneau, AK 99802–1668. Individual copies of the amendment and the EA/RIR/IRFA may be obtained from the North Pacific Fishery Management Council, P.O. Box 103136, Anchorage, AK 99501 (telephone 907–274–4563).

FOR FURTHER INFORMATION CONTACT: Jay J.C. Ginter (Fishery Management Biologist, NMFS), 907-586-7230.

SUPPLEMENTARY INFORMATION: Domestic and foreign groundfish fisheries in the exclusive economic zone (EEZ) of the BSAI area are managed in accordance with the FMP. The FMP was developed by the Council under autority of the Magnuson Fishery Conservation and Management Act (Magnuson Act) and is implemented by regulations appearing at 50 CFR 611.93 and Part 675.

The Council solicits management proposals annually from the general public, other agencies, and staff between September and December. The Council set a deadline of December 12, 1986, for receiving proposals for inclusion in Amendment 11. At its meeting on January 21–23, 1987, the Council reviewed ten amendment proposals and selected seven for tentative inclusion in Amendment 11.

The Council's Plan Team prepared a draft EA/RIR/IRFA (dated March 11, 1987) of the seven proposals for public comment as required by the National Environmental Policy Act of 1969, Executive Order 12291, and NOAA policy. The Council reviewed these documents at its March 18-20, 1987, meeting and decided to release for public comment the draft EA/RIR/IRFA (dated April 15, 1987) for six of the seven proposals. At its May 20-22, 1987, meeting, the Council considered public testimony and the recommendations of its Advisory Panel (AP), Scientific and Statistical Committee (SSC), and the

Plan Team. Public testimony included presentation of an industry-negotiated compromise supporting a split season apportionment of pollock to the IVP fishery which was perceived to address several controversial management issues. The Council approved three management proposals for inclusion in Amendment 11 and recommended them to the Secretary for approval and implementation. The Plan Team revised the EA/RIR/IRFA accordingly for Secretarial review (dated July 1987). If approved, this proposed rule would implement two of the three proposals. The third proposal revises the FMP definition of "acceptable biological catch" and does not require implementation by regulation.

A description of and the reasons for the two management proposals that would be implemented by this proposed rule are as follows:

1. Split Season Apportionment of JVP.

Under this management proposal, the amount of the total allowable catch (TAC) of pollock apportioned to IVP would be divided into two parts. Part one would be equivalent to 40 percent of the sum of the initial JVP for pollock plus 15 percent of the TAC for pollock. Part one would be made available to the JVP fishery for pollock during the period January 15 through April 15. Part two would be equivalent to the remaining IVP for pollock and would be available during the period April 16 through December 31. Amendment 11 and this proposed rule would split the annual apportionment of pollock to the IVP fishery into two parts for separate time periods only during the 1988 and 1989 fishing years.

Although the specific split season management proposal adopted by the Council was not among the original proposals considered by the Council at its March, 1987 meeting, it is similar to other split-season altrnatives that address priority access to vessels that fish for domestic processors (DAP) and roe-stripping issues and that are analyzed in the draft EA/RIR/IRFA distributed for public comment. This propsoal was presented to the Council at its May 1987 meeting as an industrynegotiated compromise among JVP and DAP fishing industry representatives. The compromise reconciled conflicting views concerning two aspects of managing the BSAI pollock fishery: access to the resource and risk of biological harm.

Access

The pollock fishery in the BSAI area has evolved since 1977 from an entirely foreign-harvested fishery to a predominantly U.S.-harvested fishery. In 1987, the TAC of pollock in the BSAI area was reserved for domestic fishermen for the first time, except for small incidental catch allowances to foreign fisheries for other species. However, of the two component parts of the domestic annual harvest (DAH), the volume of fish harvested by the JVP fishery exceeds the DAP fishery by a large margin. In 1986, the landed tonnage of pollock harvested by the JVP fishery was about 17 times that of the DAP fishery, or nearly 95 percent of the total 886,000 metric ton DAH.

A variety of economic conditions may account for this current disparity between the DAP and JVP pollock harvests. These conditions are believed by some to result in an economic incentive for domestic fishermen to make deliveries of pollock to the foreign floating processors involved in joint ventures instead of to domestic shore-based processors. Shore-based processors in Dutch Harbor and Akutan, Alaska, arguing a competitive disadvantage relative to the JVP pollock fishery, proposed an FMP amendment that would establish a 100-mile zone around these communities in which JVP operations would be prohibited. This proposal would have provided the DAP fishery with exclusive access to a substantial share of the pollock resource. The argument in favor of this action was based on the priority granted to domestic processors under the processor-preference amendments to the Magnuson Act. The Council's analysis of this action (draft EA/RIR/IRFA dated April 15, 1987), however, showed that it would have had a significantly negative economic impact on domestic fishermen delivering pollock to foreign processors in joint venture operations.

Biological Risk

The transition from foreign to domestic domination of the overall pollock harvest has been accompanied by a trend toward harvesting a larger portion of the pollock TAC earlier in the fishing year than was previously done by the foreign and early JVP fisheries. Prior to the advent of JVP fisheries in the BSAI area in 1980, generally less than 25 percent of the annual total pollock harvest was taken in the period January through April. In 1986, the JVP catch of pollock through April had increased to 40 percent of the pollock TAC. By May 2, 1987, 73 percent of the initial pollock TAC had been harvested by the JVP fishery.

One reason for this trend to an intensive early-year harvest is that pollock are aggregated in spawning

concentrations in February and March. Fishing aggregated populations of fish when the catch per unit of effort is highest is more profitable than fishing when populatiosn are dispersed. In addition, spawning aggregations of pollock contain valuable quantities of roe. At times, catch rates are so high and pollock roe is so valuable that it becomes physically necessary and economically feasible to retain only the roe and discard whole male and female carcasses, a practice known as roe stripping.

A second reason is that the JVP pollock fishery, like other groundfish fisheries, is prosecuted on an open acces, first-come, first-served basis. Fishermen who begin fishing as soon as possible will likely catch more than those who wait. A predictable result is a race for fish with each fisherman attempting to maximize his catch before the TAC is reached.

One management measure initially proposed for inclusion in Amendment 11 would have prohibited the practice of roe stripping in the JVP fishery. Advocates of this proposal argued that such a prohibition would slow the race for fish and reduce the wastage of pollock meat which is valuable later in the year, particularly for production of surimi. Discussions of this proposal by the Council and its advisory groups raised the additional concern that intensive harvesting during the spawning period ultimately may reduce pollock reproductive success, biomass, and harvests. Although there are no biological data that indicate a definite relationship between the spawning biomass of pollock and the amount of young fish that first become available to the fishery, such a relationship could exist under certain conditions or population sizes. Prudent management predicates a cautious policy against removal of too many spawning fish to protect the long-term health of the pollock resource.

Based on current trends, it is likely that the entire JVP harvest of pollock would occur during the first four months of 1988, almost entirely from spawning aggregations, and even more intensively in future years. The Council adopted the split-season proposal in part as a conservative measure to mitigate any adverse effects of concentrated harvesting on spawning pollock aggregations. The split-season proposal will reduce the amount of pollock harvested by the IVP fishery during the spawning season. In addition, while not preventing an efficient fishery for the highly valued pollock roe, it will provide a part of the pollock resource for meat

and surimi production after the primary spawning season. Finally, the split-season apportionment proposal will provide two years of relative protection during which the biological risk of an intensive roe-pollock fishery can be assessed.

During the two-year effective period of this proposal, the DAP pollock fishery in the BSAI area is expected to continue its development. In addition, more biological information likely will become available. The Council will be reconsidering the effectiveness of this management measure in light of these changes.

2. Definition of Prohibited Species

This proposal would change the prohibited species definition in the FMP and its implementing regulations to list those species or species groups which must be avoided while fishing for groundfish and, if caught incidentally, must be immediately returned to the sea with minimum injury. Listed species will include the traditional prohibited species of salmon, steelhead, halibut, herring, and king and Tanner crabs for domestic and foreign groundfish fisheries plus other non-groundfish species for the foreign fishery only. Retention of any of these species would not be allowed unless authorized by other applicable law. Such authorization would allow, for example, groundfish fishermen to retain halibut caught with hook-and-line gear during an open season for halibut specified by the International Pacific Halibut Commission.

The reason for this proposed change is that the original FMP anticipated other fishery management plans for king crab, Tanner crab, and Pacific herring. The prohibited species definition in the FMP specifically exempts species that are harvested under authorization by other FMPs, PMPs, or Federal regulations.

However, the anticipated FMPs for king crab, Tanner crab, and Pacific herring ultimately failed to be implemented or were subsequently withdrawn. This led the Council to question whether these species are correctly included in the prohibited species listing. The FMP does not attempt to manage fishing for nongroundfish species but does try to limit injury to these species by the groundfish fisheries. The problem is that the current definition, at best, does not clearly state this intent and, at worst, may in fact not protect species thought to be protected as prohibited species.

An example of this problem is king crab. In the prohibited species definition under section 14.2 of the FMP, an

exception is made for species "when * * * their retention by United States vessels is not prohibited under other FMPs or Federal regulations.' Section 14.4.2.A of the FMP reinforces this exception when it states that "United States vessels must minimize their incidental harvest of* * * any * * * species the fishery for which * * * is governed by another FMP * * *." Presently, there is no operative FMP for king crab or Federal regulation prohibiting the retention of king crab by domestic vessels. Hence, king crab is a species that fits the exception and is not prohibited. By this reading of the definition, literally all the species listed in the definition are not prohibited except for salmonids and Pacific halibut for which there are other FMPs or Federal regulations. Although there are other parts of the FMP that indicate prohibited species status for non-groundfish species, the current prohibited species definition is at fault for not clearly stating this intent.

In summary, the current definition of prohibited species in the FMP is flawed. As a result, regulations implementing the FMP, pertaining to prohibited species, contain confusing and imprecise language that may not be legally enforceable against every vessel fishing for groundfish in the EEZ off Alaska. This is especially true for Tanner and king crab species since anticipated FMPs for these species are not now in effect. This problem extends also to other non-groundfish species for which other applicable law does not exist. The proposed management measure would correct this problem.

Proposed Regulatory Amendment

NOAA proposed to make a change to the regulations implementing the FMP in addition to those proposed by the Council. This change would not implement Amendment 11, but is a modification under existing authority in the FMP. A description of and reason for this change follows.

Definition of Directed Fishing

Under Amendment 10 to the FMP, a definition of directed fishing was added to the regulations governing foreign fisheries at § 611.93(b)(1)(iii). The intention of this definition (proposed at 51 FR 45349, December 18, 1986; made final at 52 FR 8592) was to enable enforcement of directed fishing prohibitions after a prohibited species catch limit had been reached. In addition, NOAA intended that the definition of directed fishing governing foreign fisheries be consistent with that governing domestic fisheries. However,

the first occurrence of the phrase "20 percent or more of the catch, take, or harvest or to" was inadvertantly omitted from both the proposed and final rules for Amendment 10. Hence, the definition in the foreign fishery regulations is proposed to be changed to indicate that this 20 percent or more of the catch, take, or harvest at any time also will be considered in determining whether directed fishing in occurring, this change would make the BSAI area foreign fishery regulations consistent with the domestic fishery regulations pertaining to the BSAI area and the Gulf of Alaska.

Classification

This proposed rule is published under section 304(a)(1)(C)(ii) of the Magnuson Act, as amended by Pub. L. 99-659, which requires the Secretary to publish regulations proposed by a Council within 15 days of receipt of the amendment and regulations. At this time the Secretary has not determined that the amendment these regulations would implement is consistent with the national standards, other provisions of the Magnuson Act, and other applicable law. The Secretary, in making these determinations, will take into account the data and comments received during

the comment period.

The Council prepared and environmental assessment (EA) for this amendment and concluded that there will be no significant impact on the environment as a result of this rule. A copy of the EA may be obtained from the Council at the address above and comments on it are requested. The change in the definition of directed fishing is intended to enhance compliance with directed fishing prohibitions by indicating that the catch, take, or harvest at any time will be considered, in addition to the amount of fish on board, in determining whether directed fishing is occurring. this change will not affect the amount of groundfish harvested, the species taken or the location of fishing activity. As such, the Assistant Administrator has determined that this change is categorically excluded from the requirement to prepare an environmental document, as provided by NOAA Directive 02-10.

The Administrator of NOAA determined that this proposed rule is not a "major rule" requiring a regulatory impact analysis under Executive Order 12291. This determination is based on the regulatory impact review/initial regulatory flexibility analysis (RIR/ IRFA) prepared by the Council. A copy of the RIR/IRFA may be obtained from the Council at the address above.

The Council prepared an initial regulatory flexibility analysis as part of the regulatory impact review of this proposed rule. The Administrator of NOAA concludes that this proposed rule, if adopted, would have significant effects on small entities, these effects have been discussed earlier in this document relative to each specific action and in the RIR/IRFA. A copy of this analysis may be obtained from the Council at the address listed above.

This rule contains no collection-ofinformation requirement subject to the Paperwork Reduction Act.

The Council determined that this rule will be implemented in a manner that is consistent to the maximum extent practicable with the approved coastal zone management program of Alaska. This determination has been submitted for review by the responsible State agencies under section 307 of the Coastal Zone Management Act.

List of Subjects

50 CFR Part 611

Fisheries, Foreign relations, Reporting and recordkeeping requirements.

50 CFR Part 675

Fisheries, Reporting and recordkeeping requirements.

Dated: August 26, 1987.

James E. Douglas, Jr.,

Deputy Assistant Administrator For Fisheries, National Marine Fisheries Service.

For reasons set out in the preamble, 50 CFR Parts 611 and 675 are proposed to be amended as follows:

PART 611-[AMENDED]

1. The authority citation for Part 611 continues to read as follows:

Authority: 16 U.S.C. 1801 et seq., 16 U.S.C. 971 et seq., 22 U.S.C. 1971 et seq., and 16 U.S.C. 1361 et seq.

2. Section 611.93 is amended by removing paragraph (b)(1)(ii)(E) and revising paragraphs (b)(1)(ii) introductory text, (b)(1)(ii)(A), and (b)(1)(iii) to read as follows:

§ 611.93 Bering Sea and Aleutian Islands groundfish fishery. * *

(b) * * *

(ii) Categories of species. Four categories of species are recognized for regulatory purposes and they are set forth in Table 1. The term "groundfish" means species in all categories except the "prohibited species" category.

(A) The term "prohibited species" means for purposes of this section: Pacific herring (Clupea harengus pallasi); salmonids (Salmonidae);

Pacific halibut (Hippoglossus stenolepis); king crab (Paralithodes spp. and Lithodes spp.); Tanner crab (Chionoecetes spp.). Except to the extent that their harvest is authorized under other applicable law, the catch or receipt of these species must be minimized and, if caught or received, they must be returned to the sea immediately in accordance with § 611.11 of this part. Records must be maintained as required by §§ 611.9, 611.90(e)(2), and this section. Any species of fish for which there is no foreign allocation must be treated in the same manner as 'prohibited species" and records must be maintained of any catches or receipts of these species, except for "nonspecified species". Catches or receipts of "non-specified species" must be treated in the same manner as "prohibited species" but records are not required of catches or receipts of these species.

(iii) Directed fishing, with respect to any species, stock or other aggregation of fish, means fishing that is intended or can reasonably be expected to result in the catching, taking, or harvesting of quantities of such fish that amount to 20 percent or more of the catch, take, or harvest, or to 20 percent or more of the total amount of fish or fish products on board at any time. It will be a rebuttable presumption that, when any species, stock, or other aggregation of fish comprises 20 percent or more of the catch, take, or harvest, or 20 percent or more of the total amount of fish or fish products on board at any time, such fishing was directed fishing for such fish.

3. In § 611.93(B0(1)(ii) Table 1 is amended by changing the column heading "Unallocated Species" to "Prohibited Species"; revising the list of species in the column to read: "Pacific halibut, Pacific herring, salmonids, king crab, Tanner crab, and other species for which there is no allocation, except nonspecified species"; removing the column headed by "Groundfish"; and revising footnote 4 to read as follows:

Table 1.* * *

4 Must be treated in the same manner as 'prohibited species' but no records are required.

PART 675—[AMENDED]

5. The authority citation for 50 CFR Part 675 continues to read as follows:

Authority: 16 U.S.C. 1801 et seq.

6. Section 675.20 is amended by revising the heading of paragraph (b). adding a new paragraph (b)(3), and revising paragraph (c)(1) to read as follows:

§ 675.20 General limitations.

. .

(b) Apportioning the reserve, surplus DAH, and JVP.

(3) Seasonal apportionment of JVP pollock. The initial amount of pollock apportioned to JVP for each subarea in accordance with paragraph (a)(4) of this section will be divided into two parts.

(i) Part One will be 40 percent of the following sum: initial JVP plus 15 percent of the TAC for pollock. The JVP pollock harvest during the first period (defined in paragraph (b)(3)(iii) of this section) resulting from directed fishing and bycatch in fisheries for other groundfish species will be counted against Part One. When the Regional Director determines that the unharvested amount of Part One is necessary for bycatch in JVP fisheries for other groundfish species during the

first period, the Secretary will publish a notice in the Federal Register prohibiting directed fishing for pollock for the remainder of the first period. Any amount of pollock in addition to Part One necessary for bycatch in JVP fisheries for other groundfish species during the first period will be counted against Part Two.

(ii) Part Two will be any unharvested portion of Part One plus the pollock JVP remaining after the first period and as adjusted by reapportionments from reserve and DAP in accordance with paragraphs (b) (1) and (2) of this section. When the Regional Director determines that the unharvested amount of Part Two is necessary for bycatch in JVP fisheries for other groundfish species during the second period, the Secretary will publish a notice in the Federal Register prohibiting directed fishing for pollock for the remainder of the second period.

(iii) JVP pollock season. For purposes of this paragraph, the first period is that portion of the fishing year beginning

January 15 and ending April 15. The second period is that portion of the fishing year beginning April 16 and ending December 31.

(c) * * *

(1) Prohibited species, for the purpose of this part, means any of the species of Pacific salmon (Oncorhynchus spp.). steelhead trout (Salmo gairdneri or Parasalmo mykiss), Pacific halibut (Hippoglossus stenolepis), Pacific herring (Clupea harengus pallasi), king crab (Paralithodes spp. and Lithodes spp.), and Tanner crab (Chionoecetes spp.) caught by a vessel regulated under this part while fishing for groundfish in the Bering Sea and Aleutian Islands management area, unless retention is authorized by other applicable law, including the regulations of the International Pacific Halibut Commission.

[FR Doc. 87–19977 Filed 8–31–87; 8:45 am]
BILLING CODE 3510-22-M

Notices

Federal Register Vol. 52, No. 169

Tuesday, September 1, 1987

This section of the FEDERAL REGISTER contains documents other than rules or proposed rules that are applicable to the public. Notices of hearings and investigations, committee meetings, agency decisions and rulings, delegations of authority, filing of petitions and applications and agency statements of organization and functions are examples of documents appearing in this section.

Done at Washington, DC, on August 25, 1987.

J. Patrick Boyle,

Administrator, Agricultural Marketing Service.

[FR Doc. 87-20019 Filed 8-31-87; 8:45 am] BILLING CODE 3410-02-M

DEPARTMENT OF AGRICULTURE

Agricultural Marketing Service

Plant Variety Protection Advisory Board; Open Meeting

AGENCY: Agricultural Marketing Service, USDA.

ACTION: Notice of meeting.

SUMMARY: This notice sets forth the schedule and proposed agenda of a forthcoming meeting of the Plant Variety Protection Advisory Board.

DATE: Tuesday, September 22, 1987, 8:00 a.m. to 4:00 p.m., open to the public.

ADDRESS: The meeting will be held at the National Agricultural Library Building, Conference Room 1400, Beltsville, Maryland.

FOR FURTHER INFORMATION CONTACT:

Dr. Kenneth H. Evans, Executive Secretary, Plant Variety Protection Advisory Board, Room 500, National Agricultural Library Building, Beltsville, Maryland 20705 (301/344-2518).

SUPPLEMENTARY INFORMATION: Pursuant to the provisions of sec. 10(a) of the Federal Advisory Committee Act (Pub. L. 92–463), this notice is given concerning a Plant Variety Protection Advisory Board meeting. The agenda for the meeting will include discussions of: (1) The minimum difference accepted for novelty between varieties, (2) plant variety protection fees, and (3) other related topics.

The meeting is open to the public. Persons, other than members, who wish to address the Board at the meeting should contact Dr. Kenneth Evans at the above address and telephone number, prior to the meeting. Written statements may be submitted to the Board prior to or at the meeting.

(Secs. 1-145, 84 Stat. 1542, as amended (7 U.S.C. 2321 et seq.))

Federal Grain Inspection Service

Request for Designation Applicants To Provide Official Services in the Geographic Area Currently Assigned to the State of Alabama

AGENCY: Federal Grain Inspection Service (Service), USDA.

ACTION: Notice.

SUMMARY: Pursuant to the provisions of the U.S. Grain Standards Act, as Amended (Act), official agency designations shall terminate not later than triennially and may be renewed according to the criteria and procedures prescribed in the Act. This notice announces that the designation of one agency will terminate, in accordance with the Act, and requests applications from parties interested in being designated as the official agency to provide official services in the geographic area currently assigned to the specified agency. The official agency is the Alabama Department of Agriculture and Industries.

DATE: Applications to be postmarked on or before October 1, 1987.

ADDRESS: Applications must be submitted to James R. Conrad, Chief, Review Branch, Compliance Division, FGIS, USDA 1400 Independence Avenue, SW., Room 1647 South Building, Washington, DC 20250. All applications received will be made available for public inspection at this address during regular business hours.

FOR FURTHER INFORMATION CONTACT: James R. Conrad, telephone (202) 447–8525.

SUPPLEMENTARY INFORMATION: This action has been reviewed and determined not to be a rule or regulation as defined in Executive Order 12291 and Departmental Regulation 1512–1; therefore, the Executive Order and Departmental Regulation do not apply to this action.

Section 7(f)(1) of the Act specifies that

the Administrator of the Service is authorized, upon application by any qualified agency or person, to designate such agency or person to provide official services after a determination is made that the applicant is better able than any other applicant to provide official services in an assigned geographic area.

Alabama Department of Agriculture and Industries (Alabama), Beard Building, P.O. Box 3336, 1445 Federal Drive Montgomery, AL 36193, was designated under the Act as an official agency to provide inspection and weighing functions on March 1, 1985.

The official agency's designation terminates on Fedruary 28, 1988. Section 7(g)(1) of the Act states that designations of official agencies shall terminate not later than triennially and may be renewed according to the criteria and procedures prescribed in the Act.

The geographic area presently assigned to Alabama, pursuant to section 7(f)(2) of the Act, which may be assigned to the applicant selected for designation is as follows: The entire State of Alabama, except those export port locations within the State.

Interested parties, including Alabama, are hereby given opportunity to apply for official agency designation to provide the official services in the geographic area, as specified above, under the provisions of section 7(f) of the Act and § 800.196(d) of the regulations issued thereunder.

Designation in each specified geographic area is for the period beginning March 1, 1988, and ending February 28, 1991.

Parties wishing to apply for designation should contact the Review Branch, Compliance Division, at the address listed above for forms and information.

Applications and other available information will be considered in determining which applicant will be designated to provide official services in a geographic area.

(Pub. L. 94-582, 90 Stat. 2867, as amended (7 U.S.C. 71 et seq.))

Date: August 24, 1987.

J.T. Abshier,

Director, Compliance Division.
[FR Doc. 87–19892 Filed 8–31–87; 8:45 am]
BILLING CODE 3410-EN-M

Designation Renewal of the State of Oregon and Southern Illinois Agency (IL)

AGENCY: Federal Grain Inspection Service (Service); USDA.

ACTION: Notice.

SUMMARY: This notice announces the designation renewal of the Oregon Department of Agriculture (Oregon) and Southern Illinois Grain Inspection Service, Inc. (Southern Illinois), as official agencies responsible for providing official services under the U.S. Grain Standards Act, as Amended (Act).

EFFECTIVE DATE: October 1, 1987.

ADDRESS: James R. Conrad, Chief, Review Branch, Compliance Division, FGIS, USDA, 1400 Independence Avenue, SW., Room 1647 South Building, Washington, DC 20250.

FOR FURTHER INFORMATION CONTACT: James R. Conrad, telephone (202) 447– 8525.

SUPPLEMENTARY INFORMATION: This action has been reviewed and determined not to be a rule or regulation as defined in Executive Order 12291 and Departmental Regulation 1512–1; therefore, the Executive Order and Departmental Regulation do not apply to this action.

The Service announced that Oregon's and Southern Illinois' designations terminate on September 30, 1987, and requested applications for official agency designation to provide official services within specified geographic areas in the April 1, 1987, Federal Register (52 FR 10391). Applications were to be postmarked by May 1, 1987. Oregon was the only applicant for designation in its geographic area and applied for designation renewal in the area currently assigned to that agency. There were two applicants for designation in the Southern Illinois geographic area. Southern Illinois applied for designation renewal in the area currently assigned to that agency. A new corporation, Mid-America Grain Inspection Service, Inc. (Mid-America), to be located in Granite City, Illinois, was established by Scott D. Deatherage, Villa Ridge, Missouri, and Donald B. Reynolds, Rock Port, Missouri, and applied for designation in the area currently assigned to Southern Illinois.

The Service announced the applicant names in the June 1, 1987, Federal Register (52 FR 20434) and requested comments on the designation renewal of Oregon and the designation renewal of Southern Illinois or the designation of Mid-America. Comments were to be postmarked by July 16, 1987. No

comments were received regarding the designation renewal of Oregon.

A total of 17 comments were received regarding the applicants for designation in the Southern Illinois geographic area. Fourteen comments supported the designation renewal of Southern Illinois. These comments generally specified favorable qualifications concerning Southern Illinois. One of these supporting commenters referenced a boundary dispute between Southern Illinois and a neighboring official agency. This matter is discussed below. These commenters included several country elevators, grain merchandisers, farm bureaus, and an official agency.

Three commenters supported the designation of Mid-America. One commenter was a trade association. Two comments were received from an official agency, a State department of agriculture. Both of these comments supported the designation of Mid-America. One of the comments included a chronology of events concerning a boundary dispute between Southern Illinois and the State agency. Geographic areas are assigned to designated agencies pursuant to section 7(f)(2) of the Act. In addition, the Act provides that except as otherwise authorized by the Administrator official agencies can only officially inspect grain that is physically located within the geographic area assigned to the agency. The boundary at issue is a river boundary between the two agencies. The commenter concluded that based upon information submitted with its comment, it would not be advisable to renew the designation of Southern Illinois.

The Service is aware of this boundary dispute between the commenter and Southern Illinois and has been actively involved in resolving this matter in cooperation with the two agencies involved. In recent discussions with both agencies, the Service believes that a workable resolution to this matter has been achieved. The Service has and will continue to monitor the situation between the agencies but does not consider that it affects the status of either official agency at this time.

The Service evaluated all available information regarding the designation criteria in section 7(f)(1)(A) of the Act; and, in accordance with section 7(f)(1)(B), determined that Oregon is able and Southern Illinois is better able than any other applicant to provide official services in the geographic area for which the Service is renewing their designations. Effective October 1, 1987, and terminating September 30, 1990, Oregon and Southern Illinois will provide official inspection services in

their entire specified geographic area, previously described in the April 1 Federal Register.

A specified service point, for the purpose of this notice, is a city, town, or other location specified by an agency for the performance of official inspection or Class X or Class Y weighing services and where the agency and one or more of its inspectors or weighers is located. In addition to the specified service points within the assigned geographic area, an agency will provide official services not requiring an inspector or weigher to all locations within its geographic area.

Interested persons may receive a listing of an agency's specified service points by contacting either the Review Branch, Compliance Division, at the address listed above or the agencies at the following addresses: Oregon Department of Agriculture, Agriculture Building, 635 Capitol Street, NE., Salem, OR 97310–0110; and Southern Illinois Grain Inspection Service, Inc., 101 South Cherry Street, P.O. Box 630, O'Fallon, IL 62269.

[Pub. L. 94-582, 90 Stat. 2867, as amended (7 U.S.C. 71 et seq.)]

Date: August 26, 1987.

J.T. Abshier,

Director, Compliance Division.
[FR Doc. 87–19893 Filed 8–31–87; 8:45 am]
BILLING CODE 3410-EN-M

Request for Comments on Designation Applicants in the Geographic Area Currently Assigned to the Decatur Agency (IL), and State of South Carolina

AGENCY: Federal Grain Inspection Service (Service), USDA.

ACTION: Notice.

SUMMARY: This notice requests comments from interested parties on the applicants for official agency designation in the geographic area currently assigned to Decatur Grain Inspection, Inc. (Decatur), and the South Carolina Department of Agriculture (South Carolina).

DATE: Comments to be postmarked on or before October 16, 1987.

ADDRESS: Comments must be submitted in writing to Lewis Lebakken, Jr., Information Resources Staff, FGIS, USDA, Room 1661 South Building, 1400 Independence Avenue, SW., Washington, DC 20250.

Telemail users may respond to [IRSTAFF/FGIS/USDA] telemail. Telex users may respond as follows: To: Lewis Lebakken, TLX: 7607351, ANS:FGIS UC.

All comments received will be made available for public inspection at the above address during regular business hours (7 CFR 1.27(b)).

FOR FURTHER INFORMATION CONTACT: Lewis Lebakken, Jr., telephone (202) 382–1738.

SUPPLEMENTARY INFORMATION: This action has been reviewed and determined not to be a rule or regulation as defined in Executive Order 12291 and Departmental Regulation 1512–1; therefore, the Executive Order and Departmental Regulation do not apply to this action.

The Service requested applications for official agency designation to provide official services within specified geographic areas in the July 1, 1987, Federal Register (52 FR 24490). Applications were to be postmarked by July 31, 1987. Decatur and South Carolina were the only applicants for designation in their geographic area; and each applied for designation renewal in the area currently assigned to that agency.

This notice provides interested persons the opportunity to present their comments concerning the designation of the applicants. Commenters are encouraged to submit reasons for support or objection to these designation actions and include pertinent data to support their views and comments. All comments must be submitted to the Information Resources Staff, Resources Management Division, at the above address.

Comments and other available information will be considered in making a final decision. Notice of the final decision will be published in the Federal Register, and the applicants will be informed of the decision in writing.

(Pub. L. 94-582, 90 Stat. 2867, as amended (7 U.S.C. 71 et seq.))

Date: August 24, 1987.

J.T. Abshier,

Director, Compliance Division.

[FR Doc. 87-19894 Filed 8-31-87; 8:45 am] BILLING CODE 3410-EN-M

Request for Designation Applicants to Provide Official Services, in the Geographic Area Currently Assigned to Agricultural Seed, Laboratories, Inc. (AZ)

AGENCY: Federal Grain Inspection Service (Service), USDA. ACTION: Notice.

SUMMARY: This notice announces that there were no timely applicants for official agency designation in the geographic area currently assigned to Agricultural Seed Laboratories, Inc. (Agri Seed), pursuant to the July 1
Federal Register notice requesting such applicants. The Service is again requesting applications from parties interested in being designated as the official agency to provide official services in the geographic area currently assigned to Agri Seed.

DATE: Applications to be postmarked on or before October 1, 1987.

ADDRESS: Applications must be submitted to James R. Conrad, Chief, Review Branch, Compliance Division, FGIS, USDA, 1400 Independence Avenue, SW., Room 1647 South Building, Washington, DC 20250. All applications received will be made available for public inspection at this address during regular business hours.

FOR FURTHER INFORMATION CONTACT: James R. Conrad, telephone (202) 447– 8525.

SUPPLEMENTARY INFORMATION:

This action has been reviewed and determined not to be a rule or regulation as defined in Executive Order 12291 and Departmental Regulation 1512–1; therefore, the Executive Order and Departmental Regulation do not apply to this action.

Section 7(f)(1) of the Act specifies that the Administrator of the Service is authorized, upon application by any qualified agency or person, to designate such agency or person to provide official services after a determination is made that the applicant is better able than any other applicant to provide official services in an assigned geographic area.

The Service requested applications for official agency designation to provide official services within specified geographic areas in the July 1, 1987, Federal Register (52 FR 24490).

Applications were to be postmarked by July 31, 1987; we received no applications for the Agri Seed designation postmarked by that date. As a result, we are again asking for applications for designation in the Agri Seed geographic area.

Agri Seed's designation terminates on December 31, 1988. Section 7(g)(1) of the Act states that designations of official agencies shall terminate not later than triennially and may be renewed according to the criteria and procedures prescribed in the Act.

The georgraphic area presently assigned to Agri Seed, in the State of Arizona, pursuant to section 7(f)(2) of the Act, which may be assigned to the applicant selected for designation is as follows: Maricopa, Pinal, and Yuma

Interested parties, including Agri Seed, are hereby given opportunity to apply for official agency designation to provide the official services in the geographic area, as specified above, under the provisions of section 7(f) of the Act and § 800.196(d) of the regulations issued thereunder.

Designation in the specified geographic area is for the period beginning January 1, 1988, and ending December 31, 1990.

Parties wishing to apply for designation should contact the Review Branch, Compliance Division, at the address listed above for forms and information.

Applications and other available information will be considered in determining which applicant will be designated to provide official services in a geographic area.

Pub. L. 94–582, 90 Stat. 2867, as amended (7 U.S.C. 71 et seq..)

Date: August 24, 1987.

J.T. Abshier

Director, Compliance Division.
[FR Doc. 87–19895 Filed 8–31–87; 8:45 am]
BILLING CODE 3410-EN-M

Designation of Mid-Iowa Grain Inspection, Inc. (IA), In the Cedar Rapids, IA, Geographic Area

AGENCY: Federal Grain Inspection Service (Service).

ACTION: Notice.

summary: This notice announces the designation of Mid-Iowa Grain Inspection, Inc. (Mid-Iowa), as an official agency responsible for providing official services under the U.S. Grain Standards Act, as Amended (Act), in the Cedar Rapids, Iowa, geographic area.

EFFECTIVE DATE: October 1, 1987.

ADDRESS: James R. Conrad, Chief, Review Branch, Compliance Division, FGIS, USDA, 1400 Independence Avenue, SW., Room 1647 South Building, Washington, DC 20250.

FOR FURTHER INFORMATION CONTACT: James R. Conrad, telephone (202) 447–8525.

SUPPLEMENTARY INFORMATION:

This action has been reviewed and determined not to be a rule or regulation as defined in Executive Order 12291 and Departmental Regulation 1512–1; therefore, the Executive Order and Departmental Regulation do not apply to this action.

The Service announced the cancellation of designation of Cedar Rapids Grain Service, Inc., effective September 30, 1987, and requested applications for official agency designation to provide official services within a specified geographic area in the April 1, 1987 Federal Register (52 FR

10392). Applications were to be postmarked by May 1, 1987. Florian E. Polaski and Jeffrey Polaski, Cedar Rapids, Iowa, proposed to establish Mid-Iowa Grain Inspection, Inc., and applied for designation in the entire area available for assignment.

The Service announced the applicant name in the June 1, 1987, Federal Register (52 FR 20435) and requested comments on the designation of Midlowa. Comments were to be postmarked by June 15, 1987; none were received.

The Service evaluated all available information regarding the designation criteria in section 7(f)(1)(A) of the Act, and in accordance with section 7(f)(1)(B), determined that Mid-Iowa is able to provide official services in the geographic area for which the Service is designating it. Effective October 1, 1987, and terminating September 30, 1990, Mid-Iowa will provide official inspection services in the entire specified geographic area, previously described in the April 1 Federal Register.

A specified service point, for the purpose of this notice, is a city, town, or other location specified by a agency for the performance of official inspection or Class X or Class Y weighing services and where the agency and one or more of its inspectors or weighers is located. In addition to the specified service points within the assigned geograhic area, an agency will provide official services not requiring an inspector or weigher to all locations within its geographic area.

Interested persons may receive a listing of the agency's specified service points by contacting either the Review Branch, Compliance Division, at the address listed above or the agency at the following address: Mid-Iowa Grain Inspection, Inc., 1114–55th Avenue, S.W., Cedar Rapids, IA 52404.

Pub. L. 94–582, 90 Stat. 2867, as amended (7 U.S.C. 71 et seq.)

Dated: August 24, 1987.

J.T. Abshier

Director Compliance Division.

[FR Doc. 87-19896 Filed 8-31-87; 8:45 am]

COMMISSION ON CIVIL RIGHTS

Agenda and Notice of Public Meeting; California Advisory Committee

Notice is hereby given, pursuant to the provisions of the Rules and Regulations of the U.S Commission on Civil Rights, that a meeting of the California Advisory Committee to the Commission will convene at 10:00 a.m. and adjourn at 5:00 p.m., on September 11, 1987, at

Centro Maravilla, 4716 East Brooklyn Avenue, Los Angeles, California 90022. The purpose of the meeting is to conduct a community forum on the impact of the implementation of the Immigration Reform Act.

Persons desiring additional information, or planning a presentation to the Committee, should contact Committee Chairperson, Helen Hernandex or Philip Montez, Director of the Western Regional Division (213) 894–3437, (TDD 213/894–0508). Hearing impaired persons who will attend the meeting and require the services of a sign language interpreter, should contact the Regional Office at least five (5) working days before the scheduled date of the meeting.

The meeting will be conducted pursuant to the provisions of the rules and regulations of the Commission.

Susan J. Prado,

Acting Staff Director. [FR Doc. 87–20027 Filed 8–31–87; 8:45 am] BILLING CODE 6335-01-M

Agenda and Notice of Public Meeting; Hawaii Advisory Committee

Notice is hereby given, pursuant to the provisions of the Rules and Regulations of the U.S. Commission on Civil Rights, that a meeting of the Hawaii Advisory Committee to the Commission will convene at 9:00 a.m. and adjourn at 3:00 p.m., on September 16, 1987, at the Ala Moana Hotel, 410 Atkinson Drive, Honolulu, Hawaii 96814. The purpose of the meeting is to obtain information on the status of affirmative action and equal employment opportunities within the State Department of Education. A brief planning session will be convened in the afternoon.

Persons desiring additional information, or planning a presentation to the Committee, should contact Committee Chairperson, Andre S.
Tatibouet, or Philip Montez, Director of the Western Regional Division (213) 894–3437, (TDD 213/894–0508). Hearing impaired persons who will attend the meeting and require the services of a sign language interpreter, should contact the Regional Office at least five (5) working days before the scheduled date of the meeting.

The meeting will be conducted pursuant to the provisions of the rules and regulations of the Commission.

Dated at Washington, DC, August 24, 1987. Susan J. Prado,

Acting Staff Director.

[FR Doc. 87-20028 Filed 8-31-87; 8:45 am] BILLING CODE 6335-01-M

Agenda and Notice of Public Meeting; Oregon Advisory Committee

Notice is hereby given, pursuant to the provisions of the Rules and Regulations of the U.S. Commission on Civil Rights, that a meeting of the Oregon Advisory Committee to the Commission will convene at 10:00 a.m. and adjourn at 12:30 p.m., on September 18, 1987, at the Hilton Hotel, 921 Southwest 6th Avenue, Portland, Oregon 97204. The purpose of the meeting is to plan activities and programming for the coming year.

Persons desiring additional information, or planning a presentation to the Committee, should contact Committee Chairperson, James Huffman or Philip Montez, Director of the Western Regional Division (213) 894–3437, (TDD 213/894–0508). Hearing impaired persons who will attend the meeting and require the services of a sign language interpreter, should contact the Regional Office at least five (5) working days before the scheduled date of the meeting.

The meeting will be conducted pursuant to the provisions of the rules and regulations of the Commission.

Dated at Washington, DC, August 21, 1987. Susan J. Prado,

Acting Staff Director.

[FR Doc. 87-20029 Filed 8-31-87; 8:45 am] BILLING CODE 6335-01-M

DEPARTMENT OF COMMERCE

International Trade Administration

Antidumping or Countervailing Duty Order, Finding, or Suspended Investigation; Opportunity to Request Administrative Review

AGENCY: International Trade Administration/Import Administration, Department of Commerce.

ACTION: Notice of opportunity to request administrative review of antidumping or countervailing duty order, finding, or suspended investigation.

Background

Each year during the anniversary month of the publication of an antidumping or countervailing duty order, finding, or suspension of investigation, an interested party as defined in section 771(9) of the Traffic Act of 1930 may request, in accordance with § 353.53a or § 355.10 of the Commerce Regulations, that the Department of Commerce ("the Department") conduct an administratiave review of that antidumping or countervailing duty

order, finding, or suspended investigation.

Opportunity to Request a Review

Not later than September 30, 1987, interested parties may request administrative review of the following orders, findings, or suspended investigations, with anniversary dates in September for the following periods:

	Period
Antidumping Duty Proceeding	
Replacement Parts for Self-Pro- pelled Bituminous Paving Equip-	
ment from Canada	09/01/86-08/31/87
Steel Jacks from Canada	09/01/86-08/31/87
ming Pools from Japan Carbon Steel Bars and Structural	09/01/86-08/31/87
Shapes from Canada	09/01/86-08/31/87
from the People's Republic of China	
Countervalling Duty Proceeding	
Portland Hydraulic Cement and	
Cement Clinker form Mexico	01/01/86-12/31/87
Lamb Meat from New Zealand	04/01/86-03/31/87
Lime from Mexico	01/01/86-12/31/86
Fresh Cut Roses from Israel	01/01/86-09/30/86
Steel Wire from New Zealand	09/02/86-12/31/86
Carbon Steel Wire Rod from Argen-	
tina	01/01/86-12/31/88
Shop Towels from Peru	01/01/86-12/31/86

Seven copies of the request should be submitted to the Deputy Assistant Secretary for Import Administration, International Trade Administration, Room B-099, U.S. Department of Commerce, Washington, DC 20230.

The Department will publish in the Federal Register a notice of "Initiation of Antidumping (Countervailing) Duty Administrative Review," for requests received by September 30, 1987.

If the Department does not receive by September 30, 1987 a request for review of entries covered by an order or finding listed in this notice and for the period identified above, the Department will instruct the Customs Service to assess antidumping or countervailing duties on those entries at a rate to the cash deposit of (or bond for) estimated antidumping or countervailing duties required on those entries at the time of entry, or withdrawal from warehouse, for consumption and to continue to collect the cash deposit previously ordered.

This notice is not required by statute, but is published as a service to the international trading community.

Joseph A. Spetrini,

Acting Deputy Assistant Secretary For Import Administration.

Date: August 24, 1987. [FR Doc. 87–20058 Filed 8–31–87; 8:45 am] BILLING CODE 3510–DS-M

[A-412-602]

Final Determination of Sales at Less Than Fair Value, Certain Forged Steel Crankshafts From the United Kingdom

AGENCY: International Trade Administration, Import Administration, Commerce.

ACTION: Notice.

SUMMARY: We determine that certain forged steel crankshafts (CFSC) from the United Kingdom (U.K.) are being, or are likely to be, sold in the United States at less than fair value. We have notified the U.S. International Trade Commission (ITC) of our determination and have directed the U.S. Customs Service to continue to suspend liquidation of all entries of CFSC from the U.K. that are entered or withdrawn from warehouse, for consumption, on or after the date of publication of this notice, and to require a cash deposit or bond for each entry in an amount equal to the estimated weighted-average dumping margins as described in the "Suspension of Liquidation" section of

EFFECTIVE DATE: September 1, 1987.

FOR FURTHER INFORMATION CONTACT:
Ms. Loc Nguyen, Ms. Lori Cooper, or Ms.
Barbara Tillman, Office of
Investigations, Import Administration,
International Trade Administration, U.S.
Department of Commerce, 14th Street
and Constitution Avenue, NW,
Washington DC 20230; telephone: (202)
377-0167, 377-8320, or 377-2438.

SUPPLEMENTARY INFORMATION:

Final Determination

We determine that imports of CFSC from the U.K. are being, or are likely to be, sold in the United States at less than fair value, as provided in section 735(a)(2) of the Tariff Act of 1930, as amended (the Act) [19 USC 1673d(a)]. We made fair value comparisons on sales of CFSC to the United States by the respondent during the period of investigation (October 1, 1985, through October 31, 1986). The estimated weighted-average dumping margins are shown in the "Suspension of Liquidation" section of this notice.

Case History

Since the last Federal Register publication pertaining to this case [the preliminary determination of sales at less than fair value (52 FR 18000, May 13, 1987)], the following events have occurred. We conducted verification from May 13–22, and on June 11, 1987, of the questionnaire responses of United Engineering & Forging (UEF). A public hearing was held on July 16, 1987.

Petitioner and respondent filed prehearing briefs on July 13, and posthearing briefs, including comments on the verification report, on July 24, 1987.

Scope of Investigation

The products covered by this investigation are forged carbon or alloy steel crankshafts with a shipping weight between 40 and 750 pounds, whether machined or unmachined. These products are currently classified under items 660.6713, 660.6727, 660.6747, 660.7113, 660.7127, and 660.7147 of the Tariff Schedules of the United States Annotated (TSUSA). Neither cast crankshafts nor forged crankshafts with shipping weights of less than 40 pounds or greater than 750 pounds are subject to this investigation.

Period of Investigation

CFSC are normally sold to the United States on the basis of long-term requirements contracts. Therefore, in order to capture the most recent sales of CFSC to the United States, we extended the period of investigation (POI) to encompass the 13 months from October 1, 1985, to October 31, 1986, as permitted by § 353.38(a) of our regulations.

Fair Value Comparisons

To determine whether sales of CFSC in the United States were made at less than fair value, we compared the United States price to the foreign market value for the company under investigation, as specified below. We made comparisons on virtually all of the sales of CFSC to the United States during the POI.

United States Price

As provided in section 772(b) of the Act, we used the purchase price of CFSC to represent the United States price for sales by UEF, because the merchandise was sold directly to unrelated purchasers prior to its importation into the United States.

We calculated the purchase price based on the c.i.f. delivered, duty-paid price to unrelated purchasers. We made deductions, where appropriate, for foreign inland, ocean and U.S. inland freight, marine insurance, U.S. customs duties, and brokerage and handling fees.

Foreign Market Value

In accordance with section 773(a)(1)(A) of the Act, we calculated foreign market value for CFSC based on delivered prices in the home market. We made deductions for foreign inland freight. Since no packing costs were incurred in the home market, we have only added U.S. packing costs. Pursuant to § 353.15(a) of our regulations, we

made circumstances of sale adjustments for differences in warranty and credit expenses. We made an adjustment to account for differences in physical characteristics of the merchandise in accordance with § 353.16 of our

regulations.

In our preliminary determination, we made no adjustment for what respondent reported as technical services expenses, because we did not consider them to be directly related expenses within the meaning of § 353.15 of our regulations. At verification, we confirmed that these expenses were not directly related to the sales under consideration. On this basis, we have not made a circumstances of sale adjustment for these expenses.

In our preliminary determination. based on information provided in UEF's response, we made an adjustment for what we believed were after-sale warehousing expenses. During verification, we found that one shipment of crankshafts was held in UEF's rental facilities in the U.S. as buffer stock, to dampen fluctuations in shipping time and customer schedules. We also found that two other shipments of crankshafts were held in the customer's warehouse. Because the factual situation pertaining to these three transactions was not established until the verification, and because they comprise less than four percent of the total value of crankshafts sold to the United States during the POL we have not included these three transactions in our fair value comparisons.

Currency Conversion

When calculating foreign market value, we made currency conversions from British pound sterling to U.S. dollars in accordance with § 353.56(a) of our regulations, using certified exchange rates furnished by the Federal Reserve Bank of New York.

Petitioner's Comments

Comment 1: Petitioner argues that, contrary to respondent's arguments, the Department should not enlarge the POI to cover sales of certain die numbers that took place prior to October 1, 1985.

Petitioner argues that, given the prevalence of long-term contracts in this industry, it is recent "sales * * * that are the appropriate focus of DOC's inquiry."

DOC Position: We agree. We believe that the 13-month POI, October 1, 1985, through October 31, 1986, set at the beginning of this investigation captures the most recent sales, allowing us to do a meaningful analysis of this case.

Comment 2: Petitioner argues that the "date of sale" should be the "date of price determination" and not the

effective date as respondent argues. Furthermore, petitioner argues that "the date of sale" should be the date on which agreement is reached as to firm price and quantity terms and not the date of the purchase order, or the date of written confirmation of an agreement.

DOC Position: We agree that the "date of sale" is the date on which all basic terms of the sale are agreed to, including the determination of price. We believe that, in this case, the date of sale is the date the price is confirmed in writing since that is the first date the

price is finalized.

Comment 3: Petitioner argues that weight ought to be a primary criterion of similarity and that only crankshafts within a 15 percent weight range should be compared. Although this 15 percent rule is not recognized in the industry as based upon any principle of forged crankshaft production, petitioner argues that "there is an obvious need to draw the line somewhere," in order to minimize the size of the physical difference adjustments. Petitioner cites several cases to support its argument about the use of a range within which "similar" products are grouped including, among others: Color Picture Tubes from Canada, 52 FR 24316, 28317 (1987) and Certain Electric Motors from Japan, 49 FR 32627 (1984).

DOC Position: We disagree. Petitioner has not provided us with any evidence supporting a cut-off point of plus or minus 15 percent. However, we have used weight as one of the major criteria by which we determined appropriate comparisons. In the cases cited above, the products covered by those investigations are sold in specific sizes: therefore, it is appropriate to use a range of sizes within which to group similar products. Crankshafts, on the other hand, are made to each customer's specifications. Therefore, although weight is a factor in choosing the most similar merchandise, the weight range itself is not the basis for establishing categories of such or similar

merchandise.

Comment 4: Petitioner contends that UEF's argument that section 771(16) of the Act requires the Department to take into consideration both physical (such as "complexity of crankshaft design") and non-physical (such as "sales" or "planning volume") characteristics in determining product "similarity" is a misinterpretation of the statute. Petitioner admits that section 771(16) does refer to such non-physical characteristics as end-use and commercial value; however, when it comes to determining what is "most similar", the statute clearly makes physical characteristics the primary

criteria. Petitioner further argues that "twisting" is not a physical characteristic, and that the physical characteristics of the home market models used by the Department in its preliminary determination are wholly unrelated to the fact that they are produced using different manufacturing techniques. Petitioner argues that "conceptually, it would appear more appropriate to consider twisting, like production volume, as a cost issue cognizable, if at all, under the commercial value criterion of the statute and therefore of much less importance than physical characteristics such as configuration and weight."

DOC Position: We disagree. Based on the evidence produced during this proceeding, we consider twisting to be as much of a physical characteristic as configuration and weight; therefore, it is one of the primary criteria in determining "most similar" products. We agree, however, that such nonphysical characteristics as sales and planning volume are not relevant for the purpose of selecting "most similar" products. See DOC Position on Respondent's Comment 2.

Comment 5: Petitioner argues that the Department should reject respondent's argument that twisted and non-twisted crankshafts are not comparable, because: (1) Petitioner has been prejudiced by UEF's untimely submission of "voluminous arguments" in support of this change in the Department's analysis this late in the investigation; (2) these arguments are unverifiable; (3) petitioner is further prejudiced by its inability to respond fully to the highly technical arguments offered by UEF; (4) UEF was unable to provide the Department with actual cost data showing that twisted crankshafts have higher costs or higher prices because they are twisted; (5) contrary to respondent's claim that it has not calculated many of the costs that go into the making of a twisted crankshaft. these cost differences have already, in fact, been quantified and furnished to the Department; and (6) judging from photographs provided, the two twisted crankshafts involved could be produced using the forged-in-position process and, therefore, are no more "complex" in shape than the two "stepped" crankshafts shown in the photographs. Finally, petitioner argues that the additional cost of twisting is not a material factor in total manufacturing costs and that the small cost discrepancy is irrelevant to a pricing decision.

DOC Position: We disagree. The issue of twisted crankshafts versus non-

twisted crankshafts was raised early on in this investigation. Petitioner had ample opportunity to comment on this issue. Furthermore, while it is true that UEF was unable to provide actual cost data, the issue is whether a twisted crankshaft is sufficiently physically similar to a non-twisted crankshaft to allow comparison. Costs relating to physical differences are relevant only once we have determined that the crankshafts are similar. Since we determined that other, non-twisted crankshafts were more similar to nontwisted crankshafts for comparison purposes, the cost of producing a twisted crankshaft is irrelevant, as is the actual production process used. Furthermore, we verified that the crankshafts were actually "twisted" rather than "forged-in-position". Thus, we determined not to compare with nontwisted crankshafts.

Comment 6: Petitioner argues that the Department should adhere to the product comparisons made in the preliminary determination, because the home market comparisons selected by the Department were "more similar" to the U.S. crankshafts than those preferred by UEF. Should the Department conclude that UEF's choices are more similar, petitioner argues that the weighted average of all home market crankshafts with the same number of throws and falling within the 15 percent weight range should be used for comparison purposes. While petitioner believes it is reasonable to compare two crankshafts similar in configuration and weight, deciding which one of two or more home market crankshafts meeting this general description is "most similar" to the U.S. crankshaft may well be a difficult, if not an impossible, task.

DOC Position: In selecting comparable products for the preliminary determination, we took into account the criteria of number of throws, weight, and forging method. In light of the evidence produced during these proceedings, we have determined that it is appropriate to take into account the additional criterion of twisting. It is our policy to use the most similar home market product for comparison purposes and not to average a number of similar home market products. We do not find that the number of adjustments to price resulting from our selection of comparable models in this case is so large as to require resorting to an averaging technique such as that proposed by petitioner, nor is there any evidence that petitioner's proposal would lead to a more accurate comparison than the models we have chosen.

Comment 7: Petitioner argues that UEF has misconstrued and misapplied the "end-use" criterion of similarity. Section 771(16)(B) of the Act includes similar end-use as a criterion of comparability. Petitioner argues that the subject crankshafts and their proposed comparison models have the same over end-use and that UEF's argument regarding "end-use" pertains to the engines into which the crankshafts are incorporated, and not to the "end-use" of the crankshafts themselves. Petitioner further argues that even if the engines were sold into different markets, the Department should not examine marketplace dynamics in deciding whether certain crankshafts are "such or similar" to one another.

DOC Position: We agree. It is the enduse of the product under investigation itself that we consider in making "similar" merchandise selections, not the end-use of other products into which the product under investigation is incorporated. See DOC Position on Respondent's Comments 1 and 2.

Comment 8: Petitioner argues that the Department should continue to convert currencies using the daily exchange rate prevailing on the date of sale, rather than the six-month forward rate. Petitioner contends that UEF's discussion of forward exchange rates is no more than a description of how UEF allegedly deals with the exchange rate risk that is inherent in virtually all international sales by foreign companies. Petitioner argues that because of regualtory prescription and the Department's consistent practice of making currency conversion calculations on the basis of the exchange rate in effect on the date of sale, there is absolutely no risk of UEF being prejudiced in an antidumping investigation by reason of exchange rate movements after the date of sale, whether the sales contract lasts for one day or for five years. Since UEF knows the pound sterling prices of its sales in the home market, and since it knows the applicable exchange rate on the date of price agreement with the U.S. buyer, if it agrees to a price that is less than fair value, it has made a conscious decision to do so and cannot blame subsequent exchange rate movements for creating a dumping margin.

DOC Position: We converted currencies using the quarterly rates certified by the Federal Reserve in accordance with § 353.56(a) of the Commerce Regulations, except where the exchange rate on the date of sale varied from the quarterly rate by five percent or more. On the one date for which there was a change greater than

five percent, we used the actual daily rate, as required. See also DOC Position on Respondent's Comment 7.

Comment 9: Petitioner argues that the Department should reject UEF's "volatility" argument because UEF based its argument on rates appearing on one particular day at the beginning of each month, thereby making the movement in exchange rates appear more dramatic than if measured based on monthly average rates. Futhermore, since exchange rates in most quarters within the POI seemed just as volatile as exchange rates in the next quarter, it seems illogical to substitute one "volatile" rate for another "volatile" rate.

DOC Position: We find that evidence does not support a conclusion that respondent reacted within a reasonable period of time to "sustained" exchange rate changes. We also find that exchange rates in this case were not "temporary" or "volatile". For these reasons, we have used the certified Federal Reserve rate in effect on the date of each sale. See DOC Position on Respondent's Comments 7 and 8.

Comment 10: Petitioner and respondent make several arguments on issues relating to credit expense calculations and the allocation of aftersale warehousing expenses on three shipments which were warehoused in the United States.

DOC Position: As discussed in the Foreign Market Value section of the notice, these three shipments of crankshafts have not been included in our fair value comparisons. Therefore, the issues of credit expense calculation and the allocation of after-sale warehousing expenses are moot.

Respondent's Comments

Comment 1: Respondent contends that the home market models chosen as comparators by the Department in the preliminary determination improperly took into account only two criteria: Number of throws and weight. The Department should consider all relevant factors in making model selections, including non-physical differences. Respondent states that, in numerous other investigations, the Department has focused on non-physical differences in identifying such or similar merchandise where identical merchandise is not sold in the home market. Respondent further contends that section 771(16)(B) expressly directs the department to consider non-physical characteristics in selecting such or similar merchandise, including the purposes for which the merchandise is used and the commercial value of the merchandise. Respondent

contends that section 771(16)(C) covers an even broader grouping, i.e., same general class or kind" of merchandise, thereby inviting a wideranging consideration of all relevant factors. Respondent cites the following in support of its position: Malleable Cast Iron Pipe Fittings, Other than Grooved, from Brazil (Pipe Fittings), 51 FR 10897 (March 31, 1986); Carlisle Tire & Rubber Co. v. United States (Carlisle), 9 C.I.T. -, 622 F. Supp. 1071 (1985); Lightweight Polyester Filament Fabric from Japan (Polyester), 49 FR 472 (January 4, 1984); Lightweight Polyster Filament Fabric from the Republic of Korea (Polyester), 48 FR 49679 (October 27, 1983), Large Power Transformers from Japan (Power Transformers), 51 FR 21197 (June 11, 1986); and J. Pattison, Antidumping and Countervailing Duty Laws (1987) 5.05(1) and 5-26.

DOC Position: In light of evidence produced during this proceeding, we have selected comparable models based on the criteria used in arriving at the preliminary determination, namely number of throws, weight, and forging method, with the addition of twisting. We believe these criteria enable us to select merchandise meeting the statutory requirements for most similar

merchandise.

Respondent's arguments concerning commercial value and end-use have described the end-use of the machines in which the crankshafts are used, rather than the end-use of the crankshafts themselves. The cases cited by respondent may be distinguished on this basis and on the facts of the different industries involved. In the Polyester cases, the fabric industry had wellestablished designations for various types of merchandise, which reflected primarily physical characteristics of the merchandise, and which were agreed upon by experts in the field. There are no well-established designations for types of merchandise in the crankshaft industry. In Cartisle and Pipe Fittings, comparability decisions included consideration of the end-use of the products under investigation, but not of the products into which they were later incorporated. Finally, Power Transformers were found to be complex products which differed in unusual features, not necessarily obvious from a reading of specifications, for which price information was deemed necessary to assist in distinguishing the various products.

Comment 2: Respondent states that, under § 353.16, once comparison models are chosen, any remaining physical differences between products are subject to adjustment for cost

differences. On this basis, respondent argues that the most appropriate methodology in this investigation is to first match factors which affect physical and commercial comparability but which cannot be accounted for with a high degree of accuracy, and then to make adjustments to account for any differences due to any remaining, more readily quantifiable factors which do not match precisely. Specifically, respondent suggests matching nontwisted with non-twisted crankshafts. crankshafts with comparable volumes of sales, and crankshafts sold for incorporaton into engines with similar end-uses. In support of this proposed methodology respondent cites Certain Electric Motors from Japan, 45 FR 73723 (November 6, 1980) and Brass Sheet and Strip from the Republic of Korea (Korean Brass Sheet), 51 FR 40833 (November 10, 1986), in which the "Department concluded that the higher production costs associated with smaller production runs of one possible home market product disqualified that product from use as the comparatoreven though it was physically closer to the U.S. product." Instead, the Department chose as the home market comparator a product produced in similar volumes to the one sold in the U.S. Respondent argues that the cost of production (COP) of twisted crankshafts is substantially higher than the nontwisted crankshafts and that not all of these incremental costs are captured in its cost accounting system. Respondent also contends that volume and end-use have a direct impact on production cost and price but are factors which are not equalized by the adjustment process. Respondent contends that the Department's own regulations expressly recognize the relevance of volume in making price-to-price comparisons. citing § 353.14, which instructs that home market and U.S. price comparisons "usually will be made on sales of comparable quantities of the merchandise under consideration."

DOC Position: While we have determined that it is inappropriate to compare non-twisted to twisted crankshafts, since twisting does indicate a physical difference in merchandise, we do not consider end-use and volume to be factors in the selection of similar merchandise in this case. Under section 771(16) of the Act, which defines "such or similar" merchandise, end-use is a factor only when the end-use pertains to the product under investigation itself, not to the product into which it is incorporated. In this case, the subject crankshafts and the proposed comparison models have the same end-

use, i.e., incorporation into engines. Therefore, it is not appropriate to use the end-use of the engines themselves as a basis for comparison. As for volume, the regulation respondent cites, § 353.14. refers to appropriate comparisons made after the selection of similar merchandise. The definition of such or similar merchandise under section 771(16), does not specify volume as a criterion for choosing the most similar merchandise. Therefore, we have not considered volume in making our selection of most similar merchandise. In Korean Brass Sheet, the case cited by respondent, the Department first determined that two home market products were equally similar to the U.S. product. Comparisons were then made to the home market product for which the production run was closest to that of the U.S. product. In that case, volume was only considered after the similar merchandise selection had been made. See also DOC Position on Petitioner's Comments 4, 5, 6, and 7.

Comment 3: Respondent argues that the factors of end-use and volume support one of its proposed comparison models, because both its comparison choice and the U.S. model are used primarily for agricultural/industrial applications and in nearly identical quantities, whereas the comparator chosen by the Department for the preliminary determination is sold in smaller volumes into a high-priced, "niche", truck market.

DOC Position: We disagree. See DOC Position on Respondent's Comments 1 and 2 and on Petitioner's Comment 7.

Comment 4: Respondent suggests that the Department should question "widely disparate margins resulting from the use of basically similar home market models as comparators." Where the different margins are attributable to identifiable, distinguishing factors that affect the commercial value of the merchandise. respondent argues that the Department must eliminate the differences, either by quantifying them and making an adjustment or by identifying a more similar home market model.

DOC Position: "Widely disparate margins which result from the use of basically similar home market models as comparators" are not necessarily an indication of inappropriate comparisons but rather could be an indication of actual dumping margins. However, if those margins are solely attributable to identifiable, distinguishing factors, then the Department will attempt to eliminate the differences, either by quantifying them and making an adjustment or by identifying a more similar home market

model. In this case, we have selected a more similar home market model.

Comment 5: Respondent argues that the plus or minus 15 percent weight range proposed by petitioner as a basis for selecting crankshaft comparison models is arbitrary, has no technical or commercial basis, and overstates the importance of similarity in weight in the process of model selection.

DOC Position: We agree that petitioner has not provided any evidence other than conclusory statements to support the proposed weight limit of plus or minus 15 percent in making our product comparisons. Therefore, where appropriate, we have gone outside that weight range in selecting the most similar home market crankshaft for comparison purposes. See DOC Position on Petitioner's Comment

Comment 6: Should the Department adhere to its preliminary comparison model choices, respondent urges that the Department weight-average the two home market models (i.e., the Department's proposed comparator and respondent's proposed comparator). Respondent argues that this approach would reduce the distortion inherent in comparing models which differ in non-adjustable respects. Finally, respondent argues that weight-averaging would be consistent with petitioner's own preference.

DOC Position: See DOC Position on Petitioner's Comment 6.

Comment 7: Respondent argues that the Department should use the sixmonth forward exchange rates for currency conversions. It contends that it is common practice in the U.K. to hedge against the effect of exchange rate fluctuation by selling forward foreign currency receipts and that this has been actual UEF policy for several years. UEF argues that it would be perverse and unfair, if the company's "sound commercial practice were ignored in determining fair value, producing exchange rate dumping", the very result sought to be avoided when applying the antidumping laws in an economic environment characterized by volatile exchange rates. Respondent further argues that nothing is said in the Department's regulations barring the use of a forward exchange rate, citing 19 CFR 353.56(a), which merely requires that the conversion be made "as of the date of the purchase or agreement to purchase". Respondent argues that the regulation does not specify a daily, quarterly, or a forward rate, and that the Department has discretion as to which

DOC Position: Section 353.56(a) requires that currency conversions be

made "in accordance with the provisions of section 522 of the Tariff Act of 1930, as amended" (31 U.S.C. 5151), which provides that "[t]he Federal Reserve Bank of New York shall decide the buying rate" [31 U.S.C. 5151(e)]. The Tariff Act also directs that conversions be made at quarterly rates, unless the rate on any given day varies from the quarterly rate by five percent or more, in which case the actual daily rate is to be used [31 U.S.C. 5151(c), (d)]. Therefore, contrary to Respondent's contention, we are obliged to use quarterly rates absent the five percent variance provided for in the Tariff Act, or absent circumstances which would permit us to apply the "special rule" of § 353.56(b) of the regulations. Even if the "special rule" could be applied in this case, UEF has not provided sufficient evidence to support its assertion that its pricing is directly linked to, or based on, the sixmonth forward exchange rate.

Comment 8: Respondent argues that if forward exchange rates are not used, the Department should apply the lag rate, i.e., use the exchange rate prevailing in the calendar quarter preceding the sales date. Respondent contends that in previous cases such as Melamine Chemicals, Inc. v. United States (Melamine), 732 F.2d 924, 931 (Fed. Cir. 1984), the Court of Appeals has upheld the Department's application of the exchange rate prevailing in the quarter preceding the sales in question to prevent the imposition of antidumping duties resulting solely from temporary

currency fluctuations. Respondent states that it renegotiated its prices with one U.S. customer to take account of the strengthening of the pound, and that this is evidence of UEF's attempt to do what the statute wants foreign producers to do-to raise U.S. prices when the dollar weakens. In Brass Sheet and Strip from the Federal Republic of Germany (German Brass Sheet), 52 FR 822, 826 (January 9, 1987), the Department specifies two tests, one of which must be met before the Department will consider lagging the exchange rates in less than fair value (LTFV) investigations: (1) There has been a sustained change in exchange rates and respondents can show that they have acted within a reasonable period of time to adjust their prices to the change, or (2) dumping m argins are due solely to a temporary fluctuation in exchange rates. Respondent contends that it has met these tests.

DOC Position: We disagree. If exchange rates in this case are considered to have been characterized by "sustained" changes, respondent's evidence has not shown price readjustment or other reaction to such changes within a reasonable period of time as required by *Melamine*. Nor does the evidence support a finding of "temporary" exchange rate changes, so that the second test cited by respondent is inapplicable.

Comment 9: Respondent argues that U.S. interest rates should be used in determining the cost of credit for U.S. sales, because the U.S. rate would reflect the actual credit costs incurred by UEF had the company borrowed to finance its U.S. receivables. Respondent cites Certain Welded Pipe and Tube Products from Turkey (Welded Pipe), 51 FR 13044 (April 17, 1986), in which the Department calculated interest expense in the U.S. market based on the relevant U.S. rates.

DOC Position: We disagree. It is the Department's policy to use the home market interest rate to compute the respondent's credit expense for U.S. purchase price sales where, as in the present investigation, the respondent has not received any foreign financing. In Welded Pipe, U.S. sales were actually financed with short-term dollar-denominated financing, so the use of the weighted-average dollar interest rate was applied.

Comment 10: Respondent argues that the Department should calculate credit adjustments based on the interest rate prevailing on the date of each shipment, the rate UEF would have had to pay had it actually borrowed to finance its receivables.

DOC Position: We agree and have done so.

Comment 11: Respondent argues that the per diem cost of credit should be calculated on the basis of a 365-day year rather than a 360-day year.

DOC Position: We agree. We found that the bank used by the respondent based its interest calculations on 365 and not 360 days.

Comment 12: Respondent argues that shipments made after October 31, 1986, should not be included in fair value calculations since these shipments are no more relevant than sales prior to October 1, 1985, or subsequent to October 31, 1986.

DOC Position: We disagree. Because of contractual practice in this industry, there is a significant difference between "sales" and "shipments" in this case. A "sale" of the product is made at the time when a price agreement is reached. "Shipments" directly related to these "sales" are subsequently sent to the customer over a period of months or even years. In order for the "sale" to be included in the dumping calculation for purposes of this final determination, the date of sale, i.e., the date of written

confirmation of the price agreement, has to be within the POI, i.e., October 1, 1985, through October 31, 1986.
"Shipments", however, are not limited to the POI. As long as the "shipments" are pursuant to a "sale" made within the POI, they should be included in the calculation for purposes of the investigation. We agree, however, that sales prior to October 1, 1985, and subsequent to October 31, 1986, are irrelevant to the calculation of fair value since they are outside the POI.

Comment 13: Respondent argues that any future dumping order issued in this case should be limited only to those sales which were actually investigated. Since the Department excludes sales to some customers from the investigation because they occurred prior to October 1, 1985, it should exclude from the scope of any order which might ultimately be issued, UEF's shipments to those customers. Otherwise, some of UEF's customers would be burdened by the requirement to deposit duties on sales which were never investigated and which the Department has no basis whatever to assume were made at less than fair value. Since sales to customers whose imports were not investigated cannot be identified by tariff classification, respondent proposes that such models be identified by means of a certification mechanism similar to that used in other areas of customs law where the need arises because the rate of duty varies depending on the actual use of the imported merchandise. Such a procedure could be adopted here to permit imports of crankshafts by these customers without the deposit of duties based on unfounded and arbitrary assumptions, rather than findings based on facts.

DOC Position: We disagree. Respondent misunderstands the statutory scheme applicable to dumping investigations and orders. The result of a dumping investigation is an estimated margin which is to be applied to future entries. The Department has not actually made a determination that such future entries were dumped, since an investigation can only evaluate practices which have already occurred. Should respondent believe that the estimated margin provided in an order does not accurately reflect the actual dumping margin for future entries, its remedy is to request a review under section 751 of the Act and § 353.53(a) of our regulations.

Comment 14: Respondent argues that a circumstances of sale adjustment must be made for the cost of tooling for one die number for which the U.S. customer paid part of the cost of tooling, since the U.K. customer for the comparison model did not pay for tooling. UEF submits that the different treatment of tooling costs in the two markets warrants an adjustment for different circumstances of sale. Respondent cites as an example, Certain Forged Steel Crankshafts from the Federal Republic of Germany (German Crankshafts), 52 FR 18002, 18003 (1987).

DOC Position: Since this issue was first brought up after verification, we were unable to verify either the cost paid by the U.S. customer, or the fact that the U.K. customer did not pay for tooling for the comparison model. We therefore determined not to make a circumstances of sale adjustment. In German Crankshafts, the Department did not make a circumstances of sale adjustment between the home and the U.S. markets. Rather, we found that there was insufficient information on the home market side for us to consider an adjustment. Therefore, we did not make a determination on whether a circumstances of sale adjustment for tooling costs was appropriate.

Comment 15: Respondent states that it agrees with petitioner's argument that differences in the costs of inspection should be disregarded in calculating the difference in merchandise adjustment because these costs are not related to physical differences in merchandise.

DOC Position: We disagree. Since each crankshaft requires a different type and a different level of inspection, we consider inspection costs to be variable costs directly related to the differences in the physical characteristics of the merchandise. Therefore, these costs have been included in the calculation of the difference in merchandise adjustment.

Verification: We verified all information used in making our final determination in accordance with section 776(a) of the Act and followed standard verification procedures, including examination of relevant sales and financial records of the company under investigation.

Suspension of Liquidation: In accordance with section 733(d) of the Act, we are directing the U.S. Customs Service to continue to suspend liquidation of all entries of CFSC from the U.K. that are entered, or withdrawn from warehouse, for consumption, on or after the date of publication of this notice in the Federal Register. The U.S. Customs Service shall require a cash deposit or the posting of a bond equal to the estimated weighted-average amount

by which the foreign market value of CFSC from the U.K. exceeds the United States price, as shown in the table below. The cash deposit or bonding rate established in the preliminary determination shall remain in effect with respect to entries or withdrawals from warehouse made prior to the date of publication of this notice in the Federal Register. This suspension of liquidation will remain in effect until further notice.

Manufacturer/producer/exporter	Weighted- average margin percentage
United Engineering & Forging	14.67
All others.	14.67

ITC Notification

In accordance with section 735(d) of the Act, we have notified the ITC of our determination. In addition, we are making available to the ITC all nonprivileged and nonproprietary information relating to this investigation. We will allow the ITC access to all privileged and business proprietary information in our files. provided the ITC confirms that it will not disclose such information, either publicly or under administrative protective order, without the written consent of the Deputy Assistant Secretary for Import Administration. The ITC will determine whether these imports materially injure, or threaten material injury to, a U.S. industry within 45 days of the publication of this notice.

If the ITC determines that material injury or threat of material injury does not exist, this proceeding will be terminated and all securities posted as a result of the suspension of liquidation will be refunded or cancelled. However. if the ITC determines that such injury does exist, we will issue an antidumping duty order directing the U.S. Customs Service to assess an antidumping duty on CSFC from the U.K., entered or withdrawn from warehouse, for consumption on or after the suspension of liquidation, equal to the amount by which the foreign market value exceeds the United States price.

This determination is published pursuant to section 735(d) of the Act [19 U.S.C. 1673d(d)].

Dated: August 26, 1987.

Paul Freedenberg,

Assistant Secretary for Trade Administration. [FR Doc. 87–20056 Filed 8–31–87; 8:45 am] BILLING CODE 3510-DS-M

[A-122-006]

Final Results of Antidumping Duty Administrative Review; Steel Jacks From Canada

AGENCY: International Trade Administration/Import Administration Department of Commerce.

ACTION: Notice of final results of antidumping duty administrative review.

On September 23, 1986, the
Department of Commerce published the
preliminary results of its administrative
review of the antidumping finding on
steel jacks from Canada. The review
covers the only known manufacturer of
this merchandise covered by the finding,
J.C. Hallman Manufacturing Co., Ltd.,
and the period September 1, 1983
through August 31, 1985.

We gave interested parties an opportunity to comment on our preliminary results. We received comments from the petitioner and the respondent. Based on our analysis of the comments received, the final results of review are unchanged from those presented in the preliminary results.

EFFECTIVE DATE: September 1, 1987.

FOR FURTHER INFORMATION CONTACT: Barbara Victor or David P. Mueller, Office of Compliance, International Trade Administration, U.S. Department of Commerce, Washington, DC 20230; telephone: (202) 377-5222/2923.

SUPPLEMENTARY INFORMATION:

Background

On September 23, 1986, the Department of Commerce ("the Department") published in the Federal Register (51 FR 33795) the preliminary results of its administrative review of the antidumping finding on steel jacks from Canada (31 FR 11974, September 13, 1966). We began this review under our old regulations. After the promulgation of our new regulations, the petitioner and the respondent requested in accordance with § 353.53a(a) of the Commerce Regulations that we complete the administrative review. We have now completed the administrative review in accordance with section 751 of the Tariff Act of 1930 ("the Tariff Act").

Scope of the Review

The United States has developed a system of tariff classification based on the international harmonized system of Customs nomenclature. Congress is considering legislation to convert the United States to this Harmonized System ("HS") by January 1, 1988. In view of this, we will be providing both the appropriate Tariff Schedule of the United States ("TSUS") item numbers and the appropriate HS item numbers

with our product descriptions on a test basis, pending Congressional approval.

As with the TSUS, the HS item numbers are provided for convenience and Customs purposes. The written description remains dispositive.

We are requesting petitioners to include the appropriate HS item numbers as well as the TSUS item numbers in all new petitions filed with the Department. A reference copy of the proposed Harmonized System schedule is available for consultations in the Central Records Unit, Room B-099, U.S. Department of Commerce, 14th Street and Constitution Avenue, NW., Washington, DC 20230. Additionally, all Customs offices have reference copies, and petitioners may contact the Import Specialist at their local Customs office to consult the schedule.

Imports covered by the review are shipments of assembled and unassembled steel jacks, semi-assembled jacks, disassembled jacks including jack parts from Canada, currently classifiable under item numbers 664.1057 and 664.1081 of the Tariff Schedules of the United States Annotated. These products are currently classifiable under HS item number 8425.49.00. The review covers the only known manufacturer of this merchandise to the United States covered by the finding, and the period September 1, 1983 through August 31, 1985.

Analysis of Comments Received

We invited interested parties to comment on the preliminary results. We received comments from the petitioner and the respondent.

Comment 1: Hallman argues that it has presented a compelling reason for Commerce to accept the questionnaire response filed after the preliminary results of review, because Hallman changed counsel after the questionnaire was mailed by the Department. Hallman asserts that Commerce has the authority to accept its late response. Hallman maintains that the Commerce Department is authorized to, and must accept its late response to avoid a manifest injustice. The information contained in that response should be considered in the final results of review.

Department's Position: We disagree. The preliminary results of review were based on the best information available since Hallman did not provide a timely response to our questionnaire which was transmitted to Hallman on April 31, 1986. The Department sent Hallman a follow-up letter on June 20, 1986 stating that unless it responded within an additional fifteen days, the Department

would use the best information available.

The fact that Hallman changed counsel on September 5, 1986, long after the due date to respond to the questionnaire, is no excuse for untimely filing of information. Section 353.46 of the Commerce Regulations states that "Except in situations where it would be manifestly unjust, any information or written views submitted in connection with a proceeding shall be considered only if received within the time established by these regulations or by specific instructions applicable to any request for information; and information or written views received after such time shall not be considered in the proceeding." In this case, the Department's refusal to consider the response which was submitted long after the due date and after the publication of the preliminary results is not manifestly unjust. Moreover, it would be unjust to accept the response so late in the course of the proceeding and thereby deprive the other parties of an effective opportunity to comment.

Comment 2: Bloomfield Manufacturing Co., Inc., the petitioner, contends that the final results of review must be made on the basis of best information otherwise available. Further, the petitioner argues that the rate established in a previous review should be considered as a minimum. The petitioner has submitted constructed value data which it contends the Department should use as best information otherwise available.

Department's Position: As stated in our response to Comment 1, we agree that use of best information available is appropriate in this case. Consistent with Department policy, we determined that a previous rate constitutes the best information available in this case.

Selection of the best information available is made on a case-by-case basis. It is our policy to evaluate the nature of the information available and the degree of cooperation received in exercising our discretion to choose the appropriate information to use in such situations. We did not select the most adverse of all available alternate sources of information. We believe that the previous rate selected is both sufficient to ensure timely submissions in future administrative reviews and is in compliance with the Department's regulations and established policy.

Comment 3: Bloomfield contends that this finding should not be limited to steel jacks manufactured by Hallman only but should be considered a country-wide finding according to the current practice of the Department, which is consistent with this order as amended by the Treasury Department.

Bloomfield argues that when administered by the Treasury Department, the issuance of an antidumping finding entailed amending the Customs Regulations to include the latest finding. In this case the Customs Regulations were amended to include "steel jacks from Canada" without reference to Hallman exclusively. Bloomfield contends that since Hallman was mentioned only in the preambular language of the amendment to the Customs Regulations and not in the body of the amended language, the finding of dumping is a country-wide finding

Department's Position: We agree. The Department has reviewed the petition and all notices published during the investigatory period with respect to steel jacks from Canada. The original petition and all notices describe the scope of the investigation to include steel jacks from Canada manufactured by J.C. Hallman Manufacturing Co., Ltd., the only known manufacturer at that time. The Department has no evidence to indicate that the Treasury Department's intent was to limit this finding of dumping to J.C. Hallman Manufacturing Co., Ltd. to the exclusion of other exporters of the subject merchandise from Canada. The reference to Hallman as a sole known manufacturer of the subject merchandise is, therefore, descriptive rather than exclusionary in nature. We will instruct the Customs Service to collect the estimated duty from all other

Comment 4: Bloomfield contends that the Department should clarify the scope to include assembled steel jacks, semi-assembled jacks, disassembled jacks (kits) and jack parts. Bloomfield contends that the administration of this finding is subject to circumvention, unless the Department clarifies these parts to be specifically covered by the

scope of the finding.

Department's Position: Since this order already covers steel jacks (assembled or unassembled) (50 FR 42577, October 21, 1985), the only issue before us is whether the order also includes jack parts. We agree that separately imported parts of steel jacks are properly included in the scope of the subject order when these parts, taken together, constitute the complete jack. This determination is based on a consideration of the following criteria: (1) General physical characteristics. (2) the expectations of the ultimate purchasers, (3) the channels of trade in which the product is sold, (4) the manner in which the product is advertised and

displayed, and (5) the ultimate use of the merchandise in question. The Court of International Trade has endorsed these criteria as the appropriate ones to use in determining whether a product is within the "class or kind" of merchandise described in a prior antidumping finding. See, Kyowa Gas Chemical Industry Co., Ltd. v. United States, 582 F. Supp. 887 (C.I.T. 1984); Diversified Products Corp., v. United States, 572 F. Supp. 863 (C.I.T. 1983).

The Department has learned from the Customs Service that, in addition to shipping assembled and unassembled jacks to the United States, Hallman is also shipping jack parts separately to its subsidiary, J.C. Hallman, Inc. These parts include: large runner, small runner. steel standard, steel handle, top clamp clevis, handle socket, pitman, reversing switch and latch, and base plate. These parts are fully manufactured prior to importation, and require only painting, assembly, labeling, and deburring operations after being received by J.C. Hallman, Inc. in the United States. These are relatively minor finishing operations which do not add substantial value.

These parts are apparently not separately sold to unrelated purchasers, and there is no indication that such jack parts have any channels of trade or use other than for assembly into steel jacks. Indeed, these jack parts are not sold to unrelated purchasers until they are assembled into complete steel jacks. Since there is no separate channel of trade for jack parts we determine that the third criterion noted above is met. Similarly, we determine that both the ultimate use and the ultimate purchasers of jack parts and complete jacks are the same, because jack parts are not used in any other device. Thus, the second and the fifth criteria outlined above are met. The fourth criterion is also met because, since there is no separate channel of trade for jack parts, the only respect in which they are advertised and displayed is in the form of complete steel jacks.

Finally, with respect to the first criterion, the Department does not believe that the fact that jack parts have, in some respect, different physical characteristics from complete jacks should be controlling in this instance. They only difference between the two is that complete steel jacks are, essentially, assembled jack parts. Since the parts are imported by a related subsidiary for the sole purpose of being assembled to become complete steel jacks, these differences become nonexistent as soon as the parts are assembled.

Furthermore, the Department has broad authority to ensure that domestic industries receive the protection that our antidumping duty orders are intended to provide. An important component of this broad enforcement authority is vigorous monitoring of compliance with antidumping duty orders under section 751 of the Tariff Act. Our responsibility to enforce antidumping duty orders includes the requirement to ensure that those orders are not circumvented through their narrow and mechanical interpretation without considering the intended purpose and substantive reach of these orders.

The Department may not allow for separate importation of jack parts which, after minor finishing operations, are sold in the United States as complete jacks in circumvention of the order. Such a limited interpretation of the scope of the order would be in violation of the Department's enforcement responsibility under the antidumping laws.

Therefore, we conclude that steel jack parts are the same "class or kind" of merchandise as steel jacks and are properly included in the scope of the order when, shortly after importation, these parts are assembled to constitute

complete steel jacks.

Final Results of Review

Based on our analysis of the comments received, the final results of review are the same as those presented in the preliminary results of review and we determine that the following margin exists for the period September 1, 1983 through August 31, 1985:

Manufacturer/exporter	Margin (percent)	
J.C. Hailman Manufacturing Co., Ltd.	23.35	

The Department will instruct the Customs Service to assess antidumping duties on all appropriate entries. The Department will issue appraisement instructions directly to the Customs Service.

Further, as provided in section 751(a)(1) of the Tariff Act, the Department will instruct the Customs Service to collect a cash deposit of estimated antidumping duties based upon the above margin.

For any future entries of this merchandise from a new exporter, not covered in this or prior administrative reviews, whose first shipments occurred after August 31, 1985, and who is unrelated to any reviewed firm, or any previously reviewed firm, a cash deposit of 28.35 percent shall be required.

This deposit requirement is effective for all shipments of Canadian steel jacks or parts of steel jacks, entered or withdrawn from warehouse, for consumption on or after the date of publication of this notice and shall remain in effect until publication of the final results of the next administrative review.

This administration review and notice are in accordance with section 751(a)(1) of the Tariff Act (19 U.S.C. 1675(a)(1)) and § 353.53a of the Commerce Regulations (19 CFR 353.53a).

Date: August 24, 1987.

Joseph A. Spetrini,

Acting Deputy Assistant Secretary for Import Administration.

[FR Doc. 87-20057 Filed 8-31-87; 8:45 am] BILLING CODE 3510-DS-M

National Oceanic and Atmospheric Administration

Public Meeting; Mid-Atlantic/New England Fishery Management Councils

AGENCY: National Marine Fisheries Service, NOAA, Commerce.

The Mid-Atlantic and New England Fishery Management Councils will convene a joint public meeting, September 16, 1987, at 9 a.m., at the Holiday Inn 1776, U.S. 60 Bypass Road, Williamsburg, VA (telephone: 804-220-1776), for the purpose of holding panel discussions on joint ventures, the Americanization of the Atlantic mackerel fishery, cost effective law enforcement, and ecosystem management. The public meeting may be lengthened or shortened depending upon progress of the agenda. The Councils may go into closed session (not open to the public) to discuss personnel and/or national security matters.

FOR FURTHER INFORMATION CONTACT: John C. Bryson, Executive Driector, Mid-Atlantic Fishery Management Council, Federal Building, 300 South New Street, Room 2115, Dover, DE 19901; telephone: (302) 674–2331.

Date: August 27, 1987. Bill A. Powell,

Executive Director, National Marine Fisheries Service.

[FR Doc. 87-20044 Filed 8-31-87; 8:45 a.m.]

Public Meetings; North Pacific Fishery Management Council

AGENCY: National Marine Fisheries Service, NOAA, Commerce.

The North Pacific Fishery Management Council has scheduled separate public meetings for its Gulf of Alaska Groundfish Plan Team and the Bering Sea/Aleutian Islands Groundfish Plan Team. Both Plan Teams will be discussing development of resource assessment documents, deriving estimates of acceptable biological catch for commercial groundfish species, and reviewing public proposals for management of the sablefish fishery.

The public meetings will be held at the National Marine Fisheries Service, Northwest and Alaska Fisheries Center, 7600 Sand Point Way, NE., Building 4, Seattle, WA. The Gulf of Alaska Plan Team will convene September 8 at 9 a.m. and proceed as necessary through the following day. The Bering Sea/Aleutian Islands Plan Team will convene September 10 at 9 a.m. and continue as necessary through September 11.

FOR FURTHER INFORMATION CONTACT: The North Pacific Fishery Management Council, P.O. Box 103136, Anchorage, AK 99510; telephone: (907) 274—4563.

Date: August 27, 1987.

Bill A. Powell,

Executive Director, National Marine Fisheries Service.

[FR Doc. 87-20045 Filed 8-31-87; 8:45 am] BILLING CODE 3510-22-M

Public Meetings North Pacific Fishery Management Council

AGENCY: National Marine Fisheries Service, NOAA, Commerce.

The North Pacific Fishery Management Council will meet September 23-25, 1987, at the Hilton Hotel in Anchorage, AK. Council members will elect officers for the coming year. They also will receive status of stocks reports for groundfish in the Gulf of Alaska and the Bering Sea/ Aleutian Islands, and make initial apportionments for domestic and foreign fisheries for 1988. Final approval of harvest levels will be made in December. For Gulf of Alaska pollock, the Council will review U.S. processor needs for 1987 to determine if surpluses exist for reapportionment to the joint venture industry.

The Council, also will consider final approval of Amendment 16 to the Gulf of Alaska Groundfish Fishery Management Plan (FMP), review a draft FMP for king and Tanner crab in the Bering Sea/Aleutian Islands, and review a revised Salmon FMP.

Industry proposals for management of the sablefish fishery will be reviewed and the Council will approve an options package to go out for public review. The Council also will review and approve a halibut allocations policy and review halibut allocation proposals from industry before sending them out for public review.

The Council will hear Committee reports on the domestic observer program, joint venture permit review, by one word catch management and reporting requirement, and will review a revised draft of the Secretary's proposed uniform standards. Recommendations on two-tier fees for foreign fishing in 1988 also may be on the agenda.

The Council's Scientific and Statistical Committee and Advisory Panel will begin at 10 a.m. on September 21 at the Hilton and continue at least through September 22. The Council will begin at 9 a.m. on September 23 and continue through September 25.

Other meetings scheduled throughout the week include the Council's Crab Management Committee on the evening of September 21, the Reporting Requirements Workgroup on the evening of September 22, and the Bycatch Committee following the close of the Council's meeting on September 25. All public meetings will be held at the hotel unless otherwise announced; times and rooms will be available on the first day of Council week. Other plan team and workgroup meetings may be held on short notice during the week. The Council will meet in executive session (not open to the public) at least once during the week to review ongoing litigation and other appropriate matters.

FOR FURTHER INFORMATION CONTACT: the North Pacific Fishery Management Council, P.O. Box 103136, Anchorage, AK 99510; telephone: (907) 274–4563.

Date: August 27, 1987.

Bill A. Powell,

Executive Director, National Marine Fisheries Service.

FR Doc. 87-20046 Filed 8-31-87; 8:45 am]
BILLING CODE 3510-22-M

Public Meetings; Pacific Fishery Management Council

AGENCY: National Marine Fisheries Service, NOAA, Commerce.

The Pacific Fishery Management Council and its advisory entities will convene separate public meetings, September 14–17, 1987, at the Pony Village Lodge, Pony Village Shopping Center, North Bend, OR, as follows:

Council

On September 16 the Council will convene at 9 a.m., with a closed session (not open to the public) to discuss litigation, personnel, and other appropriate matters. At 10 a.m. the Council will commence its open session to consider administrative matters and

groundfish management. After comments from its advisory entities and the public, the Council will affirm or revise the 1987 sablefish acceptable biological catch/optimum yield, adopt groundfish management adjustments for the last trimester, consider recommendations from its Limited Entry Committee, adopt groundfish Amendment 3 for public review, adopt preliminary groundfish management specifications for 1988, conduct a scoping session for future groundfish amendments, and other groundfish matters. There will be a public comment period at 4 p.m.

On September 17 the Council will reconvene at 9 a.m. to complete any unfinished groundfish business and address halibut allocation and salmon management. After comment from its advisory entities and the public, the Council will adopt a halibut allocation process and hear a status report on initial allocation discussions. For salmon, the Council will adopt an amendment to the fishery management plan for public review, conduct a scoping session for future amendments to the fishery management plan, and consider other salmon matters.

Scientific and Statistical Committee

On September 14 will convene at 1 p.m. to consider matters on the Council's agenda, and reconvene September 15 at 8 a.m. to meet with the Groundfish Management Team to review stock assessments for 1988.

Groundfish Select Group

On September 15 will meet at 3 p.m. to formulate a recommendation to the Council on third trimester management adjustments and other matters.

Budget Committeee

On September 16 will meet at 8 a.m. to reivew the status of the budget for the remainder of the calendar year.

Detailed agendas for all of the above meetings will be available to public after August 28. For further information contact Lawrence D. Six, Executive Director, Pacific Fishery Management Council, Metro Center, 2000 SW. First Avenue, Suite 420, Portland, OR 97201; telephone: (503) 221–6352.

Date: August 27, 1987.

Bill A. Powell,

Executive Director, National Marine Fisheries Service.

[FR Doc. 87-20047 Filed 8-31-87; 8:45 am]

DEPARTMENT OF DEFENSE

Office of the Secretary

Changes in Meeting of the Defense Science Board Task Force on B-1B Defensive Avionics

ACTION: Change in date/location of advisory committee meeting notice.

SUMMARY: The meeting of the Defense Science Board Task Force on B-1B Defensive Avionics scheduled for September 17-18, 1987 as published in the Federal Register (Vol. 52, No. 129, Page 25458, Tuesday, July 7, 1987, FR Doc. 87-15410) will be held on September 16-18, 1987 at Wright Patterson AFB, Dayton, Ohio.

Patricia H. Means,

OSD Federal Register Liaison Officer, Department of Defense.

August 27, 1987.

[FR Doc. 87-20063 Filed 8-31-87; 8:45 am] BILLING CODE 3810-01-M

Cancellation of Meeting; Defense Science Board Task Force on B-1B Defensive Avionics

ACTION: Cancellation of meeting.

SUMMARY: The meeting notice for the Defense Science Board Task Force on B-1B Defensive Avionics for August 18-19, 1987 as published in the Federal Register (Vol. 52, No. 129, Page 25458, Tuesday, July 7, 1987, FR Doc. 87-15410.) has been cancelled.

August 27, 1987.

Patricia H. Means,

OSD Federal Register Liaison Officer, Department of Defense.

[FR Doc. 87-20064 Filed 8-31-87; 8:45 am]

Department of The Army

Army Science Board; Open Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), announcement is made of the following Committee Meeting:

Name of the committee: Army Science Board (ASB).

Dates of meeting: 21 and 22 September 1987.

Times of meeting: 0800-1600 hours each day.

Place: Science Applications
International Corporation McLean,
Virginia

Agenda: The ASB Ad Hoc Subgroup on U.S. Army CECOM RD&E Center Effectiveness Review will meet to review draft report material covering the review. This meeting will be open to the public. Any person may attend, appear before, or file statements with the committee at the time and in the manner permitted by the committee. The Army Science Board Administrative Officer, Sally Warner, may be contacted for further information at [202] 695–3039 or 695–7046.

Sally A. Warner,

Administrative Officer, Army Science Board. [FR Doc. 87-20030 Filed 8-31-87; 8:45 am] BILLING CODE 3710-08-M

DEPARTMENT OF EDUCATION

Invitation for Applications for New State Grant Awards for Fiscal Year 1988; Education of the Handicapped

Title program: Training Personnel for the Education of the Handicapped. CFDA no: 84.029H.

Purpose: To increase the quantity and improve the quality of personnel to educate handicapped children and youth. Applications for State grants may be submitted by State educational agencies (SEAs). SEAs that apply for a continuation grant for fiscal year 1968 are not eligible for a new State grant in fiscal year 1988.

Deadline for transmittal of applications: December 18, 1987.

Applications available: September 8, 1987.

Estimated range of awards: \$50,000-\$85,000.

Estimated average size of awards: \$75,000.

Estimated number of awards: 13. Average project period: 36 months.

Applicable regulations: (a) The Training Personnel for the Education of the Handicapped Program, 34 CFR Part 319, 52 FR 25830 et seq.; and (b) the Education Department General Administration Regulations, 34 CFR Parts 74, 75, 77, and 78.

For applications or information contact: Norman D. Howe, U.S.
Department of Education, Office of Special Education Programs, Division of Personnel Preparation, 400 Maryland Avenue, SW. (Switzer Building, Room 3094—M/S 2313), Washington, DC 20202. Telephone: (202) 732–1068.

Program authority: 20 U.S.C. 1432. (Catalog of Federal Domestic Assistance No. 84.029: Training Personnel for the Education of the Handicapped) Dated: August 26, 1987.

Madeleine Will,

Assistant Secretary, Office of Special Education.

[FR Doc. 87-20066 Filed 8-31-87; 8:45 am] BILLING CODE 4000-01-M

Invitation for Applications for New Awards for Fiscal Year 1988; Education of the Handicapped

Title of program: Training Personnel for the Education of the Handicapped. CFDA No: 84.029.

Purpose: To increase the quantity and improve the quality of personnel available to educate children and youth with handicaps.

Applications available

Applicable regulations: (a) The Training Personnel for the Education of the Handicapped Program, 34 CFR Part 318; and (b) the Education Department General Administrative Regulations, 34 CFR Parts 74, 75, 77, and 78.

Priorities

The Secretary announces, pursuant to 34 CFR 75.105(c)(3) and 318.11 the following priorities for fiscal year 1988. The Secretary will give an absolute preference to applications that meet any of the priorities.

Preparation of Special Educators (84.029B)

This priority supports projects designed to provide preservice training of personnel for careers in special education of handicapped children and youth. The priority includes the preparation of special educators of the handicapped, including personnel trained in speech, language, and hearing impairments, and adaptive physical educators.

Preparation of Leadership Personnel (84.029D)

This priority supports doctoral and post-doctoral preservice preparation of professional personnel to conduct training of teacher trainers, researchers, administrators, and other specialists.

Preparation of Personnel for Minority Handicapped Children (84.029E)

This priority supports the preservice preparation of special education and related services personnel to educate minority and underserved populations, and provides training for members of groups which have been traditionally underrepresented in these fields.

Preparation of Related Services Personnel (84.029F)

This priority supports the preservice preparation of individuals who provide developmental, corrective, and other supportive services as may be required to assist a handicapped child or youth to benefit from special education. The priority supports the preparation of paraprofessional personnel, career educators, recreation specialists, health services personnel, school psychologists, social service providers, counselors, physical therapists. occupational therapists, volunteers, and other personnel providing special services.

Preparation of Personnel for Transition of Handicapped Youth to Adult and Working Life (84.029G)

This priority supports the preservice preparation of special education and related services personnel, including secondary school teachers, who will prepare handicapped youth to meet adult roles. Personnel may be prepared to provide either short-term transitional services, or to aid in the placement of handicapped youth in long-term employment, or both. Projects supported under this priority should prepare personnel for employment in programs dsigned to prepare handicapped youth for community placement and adjustment to the community setting.

Preparation of Personnel to Work in Rural Areas (84.029])

This priority supports preservice training of personnel for rural areas. Particular attention must be given to preservice training related to the unique aspects of providing services to special populations in rural areas. Projects supported under this priority must prepare special education personnel to fill a variety of rural specific roles with handicapped students, parents, peers, and administrators.

Special Projects (84.029K)

This priority supports projects to develop and demonstrate new approaches for the preservice training purposes set forth in § 318.10(a), for preservice training of regular educators and for the inservice training of special education personnel, including classroom aides, related services personnel, and regular education personnel who serve handicapped children and youth. Project activities assisted under this priority include development, evaluation, and distribution of imaginative or innovative approaches to personnel preparation, and development of materials to prepare personnel to educate handicapped children and youth.

Parent Organization Projects (84.029M)

This priority supports grants to parent organizations as defined in 34 CFR 318.2(b), for the purpose of providing training and information to parents of handicapped children and youth, and to volunteers who work with parents to enable those individuals to participate more effectively with professionals in meeting the educational needs of handicapped children and youth.

Preparation of Personnel to Provide Special Education and Related Services to Newborn and Infant Children with Handicaps (84.029Q)

This priority supports the preservice preparation of personnel who will serve newborn and infant children with handicaps, or newborn and infant children who are determined to be at high risk of being handicapped, or both. Personnel may be prepared to provide short-term special education and related services as necessary in an intensive care nursery, or long-term special education and related services which extend into a preschool program. Projects supported under this priority prepare personnel for employment in programs characterized by strong interaction of the medical, educational, and related services communities, and by involvement of parents and guardians who are the primary care givers for their children.

For applications or information contact: Normal D. Howe, Office of Special Education Programs, U.S. Department of Education, 400 Maryland Avenue SW., (Switzer Building, Room 3094—M/S 2313), Washington, DC 20202. Telephone: (202) 732–1068.

Program authority: 20 U.S.C. 1431. (Catalog of Federal Domestic Assistance No. 84.029; Training Personnel for the Education of the Handicapped)

Dated: August 26, 1987.

Madeleine Will,

Assistant Secretary, Office of Special Education and Rehabilitative Services.

APPLICATION NOTICES FOR FISCAL YEAR 1988

Title and CFDA number	Deadline for transmittal of applications	Available funds 1	Estimated range of awards	Estimated size of awards	Estimated number of awards	Project period in months
Preparation of Special Educa-	Charles The					
tors (84.029B)	10/23/87	\$8,500,000	\$60,000-\$80,000	\$80,000	106	Up to 60
sonnel (84.029D)	11/27/87	1,000,000	70,000-90,000	90,000	11	Up to 60.
(84.029E)	04/06/88	500,000	60,000-75,000	75,000	7	Up to 60.
Personnel (84.029F) Preparation of Personnel for Transition of Handicapped Youth to Adult and Working	12/11/87	1,400,000	50,000-70,000	70,000	20	Up to 60.
Life (84.029G)	11/20/87	500,000	60,000-80,000	80,000	6	Up to 60.
(84.029J)	04/06/87	500,000	60,000-75,000	75,000	7	Up to 60.
Special Projects (84.029K)	12/18/87	900,000	65,000-85,000	85,000	10	Up to 60.
(84.029M)	12/04/87	1,000,000	80,000-110,000	110,000	9	Up to 60.
Handicaps (84.029Q)	01/08/88	750,000	60,000-75,000	75,000	10	Up to 60.

¹ The funding levels are estimated projections of available Federal resources and may be subject to revision pending changes in Congressional appropriations.

[FR Doc. 87-20067 Filed 8-31-87; 8:45 am] BILLING CODE 4000-01-M

DEPARTMENT OF ENERGY

Federal Energy Regulatory Commission

[Docket Nos. ER87-597-000 et al.]

Electric Rate and Corporate Regulation Filings; Arizona Public Service Co. et al.

August 27, 1987.

Take notice that the following filings have been made with the Commission:

1. Arizona Public Service Co.

[Docket No. ER87-597-000]

Take notice that on August 24, 1987, Arizona Public Service Company (APS) tendered for filing an Economy Energy Interchange Agreement between Arizona Public Service Company (APS) and Rocky Mountain Generation Cooperative (RMGC) executed March 30, 1987.

APS requested that this Agreement become effective 60 days from the date of filing with FERC.

This Agreement provides that Economy Energy sales by APS to RMGC shall be priced at one of the following rates: (a) A ceiling rate concept based in part on the fixed costs associated with facilities used to produce the required energy; (b) a "split-the-savings" concept; or (c) a selling price based on 120 percent of cost to produce such energy.

Copies of this filing are being served upon RMGC and the Arizona Corporation Commission.

Comment date: September 14, 1987, in accordance with Standard Paragraph E at the end of this notice.

2. Boston Edison Co.

[Docket No. ER86-405-004]

Take notice that on August 3, 1987 Boston Edison Company tendered for filing pursuant to Commission's Order dated June 18, 1987 approving the settlement agreement in these dockets, its compliance report.

Comment date: September 14, 1987, in accordance with Standard Paragraph E at the end of this notice.

Southwestern Public Service Co. and Black Mesa Power Co.

[Docket Nos. EC87-19-000 and ER87-584-000]

Take notice that on August 13, 1987, Southwestern Public Service Company (Southwestern) and Black Mesa Power Company (Black Mesa), tendered for filing a joint application seeking an order pursuant to section 203 of the Federal Power Act (FPA) and Part 33 of the Commission's regulations authorizing the sale by Southwestern of

its electric facilities located in the States of Oklahoma and Kansas to Black Mesa in exchange for approximately 626,088 shares of the Common Stock of Black Mesa (Common Stock) and cash in the amount of \$953,067.

Southwestern is an operating public utility engaged in the generation, transmission, distribution and sale of electricity for wholesale and retail uses in the States of Texas, New Mexico, Oklahoma and Kansas. Southwestern currently provides electric service in Oklahoma to communities of Beaver, Boise City, Goodwell, Guymon, Keyes, and Texhoma and surrounding areas. Southwestern also provides electric service in Kansas to the community of Elkhart and its surrounding area.

In the interest of improving administrative efficiency and reducing cost, Southwestern has formed Black Mesa, which will be a wholly-owned subsidiary upon the closing of the transaction described herein, to own and operate the electric service transmission, distribution, and general plant facilities (except for wholesale metering equipment) currently owned and operated by Southwestern in the States of Oklahoma and Kansas.

Upon consummation of the sale, Black Mesa will become a full requirements wholesale customer of Southwestern. It will continue to provide electric utility service to retail customers who reside within the affected area presently served by Southwestern.

Southwestern also tendered for filing, pursuant to section 205 of the FPA and Part 35 of the Commission's regulations, an agreement with Black Mesa for primary electric power service. The agreement provides for service under rate levels currently filed and allowed by the Commission for service by Southwestern to its existing full requirements customers.

Black Mesa has requested a determination from the Commission whether Black Mesa will be a jurisdictional utility as defined under section 201 of the FPA. If the Commission makes a determination that Black Mesa is a jurisdictional utility. Black Mesa requests, pursuant to section 205 of the FPA and Part 35 of the Commission's regulations, that the Commission accept the initial rate schedule for the provision of system access to Southwestern. The System Access Agreement to be entered into between Black Mesa and Southwestern will allow Southwestern access to Black Mesa's electric transmission and distribution facilities for the delivery of electric energy to Southwestern's wholesale customers in the States of Oklahoma, Kansas, and Texas, and for the delivery of electric energy to certain of Southwestern's retail customers located in the State of Texas.

In addition, if the Commission makes a determination that Black Mesa is a jurisdictional utility. Southwestern and Black Mesa request, pursuant to section 203 of the FPA and Part 33 of the Commission's regulations and section 204 of the FPA and Part 34 of the Commission's regulations, respectively, an order approving the acquisition by Southwestern of Black Mesa's Common Stock and the issuance by Black Mesa of its Common Stock to Southeastern, each in connection with the transaction described herein. The effective date of this transaction is expected to be September 1, 1987.

Comment date: September 14, 1987, in accordance with Standard Paragraph E at the end of this notice.

4. Montana Power Co.

[Docket No. ER87-596-000]

Take notice that on August 21, 1987, Montana Power Company (MPC) tendered for filing pursuant to section 205 of the Federal Power Act an agreement executed on July 17, 1987 (as amended) for the sale of firm energy to the Western Area Power Administration during the period from June 15, 1987 through November 30, 1987.

MPC has requested waiver of the notice provisions of § 35.3 of the Commission's regulations in order to permit the agreement to become effective as of June 15, 1987 in accordance with its terms.

Comment date: September 14, 1987, in accordance with Standard Paragraph E at the end of this notice.

5. Ogden Martin System of San Bernardino, Inc.

[Docket No. ER87-595-000]

Take notice that on August 20, 1987, Odgen Martin Systems of San Bernardino, Inc. (OMS) tendered for filing with the Federal Energy Regulatory Commission its initial rate schedule supporting documentation. The rate schedule consists of a power purchase contract between OMS and Southern California Edison Company (Edison). The power purchase contract provides for the sale of the capacity and corresponding energy of a new resource recovery and electric generating facility to be constructed in Ontario, California.

OMS has requested a waiver of notice requirements to permit filing of the rate schedule more than 120 days prior to its proposed effective date and a petition for waiver of the Commission's regulations inappropriate to qualifying small power producers including cost of service data.

Comment date: September 14, 1987, in accordance with Standard Paragraph E at the end of this notice.

6. Metropolitan Edison Co.

[Docket No. ER87-34-002]

Take notice that on July 20, 1987, Metropolitan Edison Company tendered for filing pursuant to Commission's Order dated June 18, 1987 a compliance report with refunds of excess revenue amounts and interest computed in accordance with Section 35.19a of the Commission Regulations.

The compliance report consists of: Schedule 1—Summary of refunds including interest.

Schedule 2—Monthly billing determinants and revenues including prior, present and settlement rates.

Schedule 3—Details of monthly revenue refund and associated interest.

Copies of the filing were served upon each person designated on the official service list in this proceeding.

Comment date: September 14, 1987, in accordance with Standard Paragraph E at the end of this notice.

7. Maine Yankee Atomic Power Co.

[Docket No. ER84-344-005]

Take notice that on August 14, 1987. Maine Yankee Atomic Power Company, tendered for filing pursuant to Commission's order dated July 21, 1987, two items as FPC rate schedules. The first reflects the modification of the return on common equity to 13.60%, to be effective June 1, 1987. The second contains the change in decommissioning expense, also effective June 1, 1987. That schedule further reflects proposed charges resulting from the change to decommissioning expense by wholesale customer rate group. This schedule updates Supplement No. 1 to Supplement No. 8 of Maine Yankee Rate Schedule FPC No. 1.

Copies of this filing have been served upon all parties affected by this filing.

Comment date: September 14, 1987, in accordance with Standard Paragraph E at the end of this notice.

8. The Borough of Ellwood City, Pennsylvania and Pennsylvania Power Co., Applicants

[Docket No. EC87-18-000]

Take notice that on July 30, 1987 Ellwood City Borough, Pennsylvania (Borough), and Pennsylvania Power Company (Company) tendered for filing a Joint Application under section 203 of the Federal Power Act seeking authorization for the sale of certain substation and distribution facilities by the Company to the Borough. The Ellwood City Substation facilities to be sold will enable the Borough to receive electric service from the Company at 69,000 volts rather than the current 4,160 volt level. The other facility, a distribution substation and lines, will enable the Borough to serve certain commercial customers now being served by the Company. The consideration for the facilities is \$102,734.35. A petition by the Borough for waiver from the filing fee and a petition by both parties for waiver from filing certain exhibits under § 33.3 of the Commission's Rules and Regulations were submitted with the Joint Application. The Company has also submitted an application to the Pennsylvania Public Utility Commission for approval of the transaction.

Comment date: September 14, 1987, in accordance with Standard Paragraph E at the end of this notice.

9. Central Illinois Public Service Co.

[Docket No. EL87-60-000]

Take notice that on August 24, 1987, Central Illinois Public Service Company (CIPS) tendered for filing a petition for a declaratory order disclaiming jurisdiction over a planned corporate reorganization.

CIPS is a corporation organized and existing under the laws of the State of Illinois, and is engaged in the sale of electricity which it generates, transmits, and distributes to the public in Illinois. CIPS also sells natural gas to, and transports natural gas for, the public in Illinois. CIPS owns 20 percent of the common stock of Electric Energy, Inc. (EEI), which owns a generating station at Joppa, Illinois. The remaining ownership of EEI is as follows: Union Electric Company, 40%; Illinois Power Company, 20%; and Kentucky Utilities Company 20% (together with CIPS, the "Sponsoring Companies"). EEI supplies electrical energy requirements to an installation of the Department of Energy (DOE) at Paducah, Kentucky. All of the electricity sold by EEI is sold either to the DOE or to the Sponsoring Companies. CIPS proposes to reorganize by causing the creation of a holding company (Company) which will become the owner of all the common stock of CIPS. Under CIPS' reorganization plan. CIPS will retain the whole of its facilities, as well as its ownership interest in EEL

CIPS states that the corporate reorganization does not involve any of the elements required for Federal Energy Regulatory Commission (Commission) jurisdiction under section 203 of the Federal Power Act (Act), because there will be no disposition by CIPS of its jurisdictional facilities, no merger or consolidation of jurisdictional facilities with those of another person, and no acquisition by CIPS of the securities of another public utility. CIPS further states that, even if its proposed reorganization did constitute a transaction covered by section 203, section 318 of the Act operates to deprive the Commission of jurisdiction. CIPS also states that the Commission has no jurisdiction over its proposed reorganization under section 204(a) of the Act, because no issuance of securities will be involved in the proposed reorganization. Moreover, because any issuance of securities by CIPS, in connection with the proposed reorganization, is subject to regulation by the Illinois Commerce Commission. the Commission is deprived of jurisdiction by operation of section 204(f) of the Act.

CIPS states that its proposed corporate reorganization is consistent with the public interest.

Comment date: September 14, 1987, in accordance with Standard Paragraph E at the end of this notice.

Standard Paragraph

E. Any person desiring to be heard or to protest said filing should file a motion to intervene or protest with the Federal **Energy Regulatory Commission, 825** North Capitol Street, NE., Washington, DC 20426, in accordance with Rules 211 and 214 of the Commission's Rules of Practice and Procedure (18 CFR 385.211 and 385.214). All such motions or protests should be filed on or before the comment date. Protests will be considered by the Commission in determining the appropriate action to be taken, but will not serve to make protestants parties to the proceeding. Any person wishing to become a party must file a motion to intervene. Copies of this filing are on file with the Commission and are available for public inspection.

Kenneth F. Plumb,

Secretary.

[FR Doc. 87-20080 Filed 8-31-87; 8:45 am]

FEDERAL COMMUNICATIONS COMMISSION

[MM Docket No. 86-484]

Reexamination of the Commission's Comparative Licensing, Distress Sale and Tax Certificate Policies Premised on Racial, Ethnic or Gender Classifications

AGENCY: Federal Communications Commission.

ACTION: Notice of inquiry; further extension of reply comment deadline.

SUMMARY: This action grants a motion for a further extension of time for filing reply comments in response to comments on the Notice of Inquiry in MM Docket No. 86-484 (Reexamination of the Commission's Comparative Licensing, Distress Sale and Classifications), 52 FR 596 (January 7, 1987). The initial comment filing deadline in this proceeding was originally May 7, 1987; it was later extended to June 11, 1987. The reply comment filing deadline in this proceeding was originally July 6, 1987; it was later extended to August 20, 1987. The National Black Media Coalition and other interested parties ("NBMC") filed a request for a further two-week extension of time, or until September 3, 1987. NBMC stated that a two-week extension is necessary because it is still completing its supplemental comments and legal appendices, and that the additional time will help insure that the widest possible spectrum of organizations will participate in this

proceeding. Since the Commission was persuaded that the requested extension was reasonably necessary, the Commission is again extending the reply comment date in this proceeding. Consequently, a further extension of time for filing comments until September 3, 1987, was granted.

DATES: Reply comments are now due by September 3, 1987.

ADDRESS: Federal Communications Commission, Washington, DC 20554.

FOR FURTHER INFORMATION CONTACT: Terry L. Haines, Policy and Rules Division, Mass Media Bureau, (202) 632– 7792.

SUPPLEMENTARY INFORMATION: The full text of this Commission decision is available for inspection and copying during normal business hours in the FCC Dockets Branch (Room 230), 1919 M Street NW., Washington, DC 20554. The complete text of this decision may also be purchased from the Commission's copy contractor, International Transcription Service, (202) 857–3800, 2100 M Street NW., Suite 140, Washington, DC 20037.

Federal Communications Commission.
William H. Johnson,
Acting Chief, Mass Media Bureau.
[FR Doc. 87–19937 Filed 8–31–87; 8:45 am]
BILLING CODE 6712–01–M

FEDERAL EMERGENCY MANAGEMENT AGENCY

[FEMA-798-DR]

Major Disaster and Related Determinations; Illinois

AGENCY: Federal Emergency Management Agency. ACTION: Notice.

SUMMARY: This is a notice of the Presidential declaration of a major disaster for the State of Illinois, (FEMA-798-DR), dated August 21, 1987, and related determinations.

DATED: August 21, 1987.

FOR FURTHER INFORMATION CONTACT: Sewall H. E. Johnson, Disaster Assistance Programs, Federal Emergency Management Agency, Washington, DC 20472, (202) 646–3616.

NOTICE: Notice is hereby given that, in a letter of August 21, 1987, the President declared a major disaster under the authority of the Disaster Relief Act of 1974, as amended (42 U.S.C. 5121 et seq., Pub. L. 93–288), as follows:

I have determined that the damage in certain areas of the State of Illinois resulting from severe storms and flooding beginning on August 13, 1987, is of sufficient severity and magnitude to warrant a major-disaster declaration under Public Law 93–288. I therefore declare that such a major disaster exists in the State of Illinois.

In order to provide Federal assistance, you are hereby authorized to allocate from funds available for these purposes, such amounts as you find necessary for Federal disaster assistance and administrative expenses. Consistent with the requirement that Federal assistance be supplemental, any Federal funds provided under PL 93-288 for Public Assistance will be limited to 75 percent of total eligible costs in the designated area.

Pursuant to section 408(b) of PL 93-286, you are authorized to advance to the State its 25 percent share of the Individual and Family, Grant program, to be repaid to the United States by the State when it is able to do so.

The time period prescribed for the implementation of section 313(a), priority to certain applications for public facility and public housing assistance, shall be for a period not to exceed six months after the date of this declaration.

Notice is hereby given that pursuant to the authority vested in the Director of the Federal Emergency Management Agency under Executive Order 12148, I hereby appoint Mr. Ronald Buddecke of the Federal Emergency Management Agency to act as the Federal Coordinating Officer for this declared disaster.

I do hereby determine the following areas of the State of Illinois to have been affected adversely by this declared major disaster: Elk Grove, Hanover, Leyden, Lyons, Maine, Norwood Park, Palatine, Proviso, River Forest, Riverside, Schaumburg, and Wheeling Townships in Cook County; and Addison, Bloomingdale, Downers Grove, Wayne, and York Townships in DuPage County for Individual Assistance.

(Catalog of Federal Domestic Assistance No. 83.516, Disaster Assistance.)

Julius W. Becton, Jr.,

Director.

[FR Doc. 87–20016 Filed 8–31–87; 8:45 am] BILLING CODE 6718-02-M

[FEMA REP-2, Rev. 1]

Guidance on Offsite Emergency Radiation Measurement Systems; Phase 1—Airborne Release

AGENCY: Federal Emergency Management Agency (FEMA), ACTION: Extension of due date for submittal of comments.

SUMMARY: The document, Guidance on Offsite Emergency Radiation Measurement Systems, Phase 1Airborne Release, FEMA REP-2, Rev. 1, dated July 1987, was available for public distribution and comment on July 20, 1987, as indicated in the notice of availability of FR Vol. 52, No. 117, page 23210.

The due date for the receipt of comments has been extended.
Comments on this document will be accepted through September 30, 1987, and should be addressed to: Rules Docket Clerk, Federal Emergency Management Agency, Room 835, 500 C Street Southwest, Washington, DC 20472.

Dated: August 26, 1987.

For the Federal Emergency Management Agency

Dave McLoughlin.

Deputy Associate Director, State and Local Programs and Support.

[FR Doc. 87-20017 Filed 8-31-87; 8:45 am] BILLING CODE 67:8-20-M

FEDERAL HOME LOAN BANK BOARD

[No. AC-657]

Final Action; Approval of Conversion Application; Capital Savings Bank, FSB, Olympia, WA

Date: August 25, 1987.

Notice is hereby given that on August 13, 1987, the Office of the General Counsel of the Federal Home Loan Bank Board, acting pursuant to the authority delegated to the General Counsel or his designee, approved the application of Capital Savings Bank, FSB, Olympia, Washington, for permission to convert to the stock form of organization. Copies of the application are available for inspection at the Office of the Secretariat at the Federal Home Loan Bank Board, 1700 G Street NW., Washington, DC 20552, and at the Office of the Supervisory Agent at the Federal Home Loan Bank of San Francisco, 600 California Street, San Francisco. California 94129, and at the Office of Supervisory Agent at the Federal Home Loan Bank of Seattle, 1501 4th Avenue, 19th Floor, Seattle, Washington 98101-

By the Federal Home Loan Bank Board. John F. Ghizzoni,

Assistant Secretary.

[FR Doc. 87-20037 Filed 8-31-87; 8:45 am] BILLING CODE 6720-01-M

[No. AC-656]

Final Action; Approval of Conversion Application; First Federal Savings and Loan Association of Lacrosse, Lacrosse, WI

Date: August 25, 1987.

Notice is hereby given that on August 21, 1987, the Office of the General Counsel of the Federal Home Loan Bank Board, acting pursuant to the authority delegated to the General Counsel or his designee, approved the application of First Federal Savings and Loan Association of LaCrosse, LaCrosse, Wisconsin, for permission to covnert to the stock form of organization. Copies of the application are available for inspection at the Office of the Secretariat at the Federal Home Loan Bank Board, 1700 G Street NW., Washington, DC 20552, and at the Office of the Supervisory Agent at the Federal Home Loan Bank of Chicago, 111 East Wacker Drive, Suite 800, Chicago, Illinois 60601.

By the Federal Home Loan Bank Board.
John F. Ghizzoni,
Assistant Secretary.
[FR. Doc. 87–20038 Filed 8–31–87; 8:45 am]
BILLING CODE 6720–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 87C-0253]

Filing of Color Additive Petition; CooperVision, Inc.

AGENCY: Food and Drug Administration.
ACTION: Notice.

SUMMARY: The Food and Drug
Administration (FDA) is announcing
that CooperVision, Inc., has filed a
petition proposing that the color
additive regulations be amended to
provide for the safe use of chromiumcobalt-aluminum oxide to color contact
lenses.

FOR FURTHER INFORMATION CONTACT: Mary W. Lipien, Center for Food Safety and Applied Nutrition (HFF-335), Food and Drug Administration, 200 C Street SW., Washington, DC 20204, 202-473-

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (section 706(d)(1), 74 Stat. 402 through 403 (21 U.S.C. 376(d)(1))), notice is given that a petition (CAP 7C0209) has been filed by CooperVision, Inc., 2610 Orchard Parkway, San Jose, CA 95134, Proposing that Part 73 of the color additive regulations be amended to provide for the safe use of chromiumcobalt-aluminum oxide to color contact lenses.

The potential environmental impact of this action is being reviewed. If the agency finds that an environmenal impact statement is not required and this petition results in a regulation, the notice of availability of the agency's finding of no significant impact and the evidence supporting that finding will be published with the regulation in the Federal Register in accordance with 21 CFR 25.40(c).

Dated: August 24, 1987.

Fred R. Shank,

Acting Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 87-20013 Filed 8-31-87; 8:45 am]

National Institutes of Health

Meeting of the Sickle cell Disease Advisory Committee; National Heart, Lung, and Blood Institute

Pursuant to Pub. L. 92–463, notice is hereby given of the meeting of the Sickle Cell Disease Advisory Committee, Division of Blodd Diseases and Resources, National Heart, Lung, and Blood Institute, October 2, 1987. The meeting will be held at the National Institutes of Health, Bethesda, Maryland 20892, Building 31, Conference Room 8, C-Wing.

The entire meeting will be open to the public from 9 a.m. to 5 p.m., to discuss recommendations on the implementation and evaluation of the Sickle Cell Disease Program.

Attendance by the public will be limited to space available.

Ms. Terry Bellicha, Chief, Communications and Public Information Branch National Heart, Lung, and Blood Institute, National Institutes of Health, Building 31, Room 4A21, (301) 496–4236, will provide a summary of the meeting and a roster of the committee members upon request.

Dr. Clarice D. Reid, Chief, Sickle Cell Disease Branch, Division of Blood Diseases and Resources, NHLBI, Federal Building, Room 508, (301) 496–6931, will furnish substantive program information.

(Catalog of Federal Domestic Assistance

Program No. 13.839, Blood Diseases and Resources Research, National Institutes of Health)

Dated: August 26, 1987,

Betty J. Beveridge,

Committee Management Officer, NIH. [FR Doc. 87–20051 Filed 8–31–87; 8:45 am] BILLING CODE 4140–01–M

Meeting of Pulmonary Diseases Advisory Committee; National Heart, Lung, and Blood Institute

Pursuant to Pub. L. 92–463, notice is hereby given of the meeting of the Pulmonary Diseases Advisory Committee, National Heart, Lung, and Blood Institute, October 29–30, 1987, at the National Institutes of Health, 9000 Rockville Pike, Building 31, Conference Room 8, Bethesda, Maryland 20892.

The entire meeting, from 8:30 a.m. on October 29 to adjournment on October 30, will be open to the public. The Committee will discuss the current status of the Division of Lung Diseases' programs and Committee plans for fiscal year 1988. Attendance by the public will be limited to the space available.

Terry Bellicha, Čhief, Communications and Public Information Branch, National Heart, Lung, and Blood Institute, Building 31, Room 4A–21, National Institutes of Health, Bethesda, Maryland 20892, (301) 496–4236, will provide a summary of the meeting and a roster of the committee members.

Dr. Suzanne S. Hurd, Executive Secretary of the committee, Westwood Building, Room 6A16, National Institutes of Health, Bethesda, Maryland 20892, (301) 496–7208, will furnish substantive program information.

(Catalog of Federal Domestic Assistance Program No. 13.838, Lung Diseases Research, National Institutes of Health)

Dated: August 26, 1987.

Betty J. Beveridge,

Committee Management Officer, NIH. [FR Doc. 87–20052 Filed 8–31–87; 8:45 am] BILLING CODE 4140-01-M

Public Health Service

Assessment of Medical Technology Implantable Pumps for Morphine

The Public Health Service (PHS), through the Office of Health Technology Assessment (OHTA), announces that it is coordinating an assessment of what is known of the safety, clinical effectiveness, and indications for the use of an implanted infusion pump to administer morphine or other narcotic or non-narcotic analgesics for the treatment of intractable cancer pain.

The use of oral and parenteral analgesics is established for the treatment of patients with intractable pain due to cancer. The development of implantable pumps has permitted the continuous administration of parenteral analgesics on an ambulatory basis. This assessment seeks to identify those clinical situations in which use of an implantable pump for analgesia would be the appropriate treatment for a patient with intractable pain due to cancer. The assessment further seeks to determine what is known about the advantages and risks of this mode of treatment compared to other approaches. This assessment also seeks to determine whether there are categories of patients for whom this mode of therapy is the treatment of choice.

PHS assessments consist of a synthesis of information obtained from appropriate organizations in the private sector and from PHS and other agencies in the Federal Government. PHS assessments are based on the most current knowledge concerning the safety and clinical effectiveness of a technology. Based on this assessment, a PHS recommendation will be formulated to assist the Health Care Financing Administration (HCFA) in establishing Medicare coverage policy. The information being sought is a review and assessment of past, current, and planned research related to this technology, a bibliography of published. controlled clinical trials and other welldesigned clinical studies. Information related to the characterization of the patient population most likely to benefit from it, as well as on the clinical acceptability and effectiveness of this technology and extent of use is also being sought. Proprietary information is not being sought. Any person or group wishing to provide OHTA with information relevant to this assessment should do so in writing no later than November 30, 1987.

Written material should be submitted to: Diane L. Adams, M.D., M.P.H., Office of Health Technology Assessment, 5600 Fishers Lane, Room 18A–27, Rockville, MD 20857, (301) 443–4990.

Date: August 20, 1987.

Morgan N. Jackson,

Acting Director, Office of Health Technology Assessment, National Center for Health Services Research and Health Care Technology Assessment.

[FR Doc. 87-20018 Filed 8-31-87; 8:45 am]
BILLING CODE 4160-17-M

DEPARTMENT OF THE INTERIOR

National Strategic Materials and Minerals Program Advisory Committee; Meeting

Notice is hereby given, in accordance with the Federal Advisory Committee Act, that the National Strategic Materials and Minerals Program Advisory Committee (NSMMPAC) will meet on Tuesday, September 29, 1987 from 9:30 a.m. until 12:30 p.m. or until business is concluded. The meeting will convene in Rooms 7000 A&B at the Department of the Interior, 18th & C Streets, NW., Washington, DC. It will be open to the public, subject to the availability of space.

The agenda will include: reports of task force activities, presentation of briefing on mine waste regulatory issues, and any recommendations for possible action by the Committee.

Statements are invited from groups and members of the general public who have an interest in mining, minerals or materials issues. The Committee is particularly interested in hearing any comments or suggestions regarding advanced materials and technology issues which fall within the purview of the Committee. To ensure that time will be available to hear such statements, prospective witnesses are requested to notify the Executive Director (see below) of their intention to appear.

FOR FURTHER INFORMATION CONTACT: Gully Walter, Department of the Interior, Washington, DC, Room 6650 (202) 343–2136.

Dated: August 27, 1987.

Gully Walter,

Executive Director.

[FR Doc. 87-20059 Filed 8-31-87; 8:45 am]

BILLING CODE 4310-10-M

Bureau of Indian Affairs

Information Collection Submitted to the Office of Management and Budget for Review Under the Paperwork Reduction Act

The proposal for the collection of information listed below has been submitted to the Office of Management and Budget for approval under the provision of the Paperwork Reduction Act (44 U.S.C. Chapter 35). Copies of the proposed information collection and related forms and explantory material may be obtained by contacting the Bureau's clearance officer at the phone number listed below. Comments and suggestions on the requirement should be made within 30 days directly to the Bureau clearance officer and to the

Office of Management and Budget Interior Department Desk Officer, Washington, DC 20503, telephone (202) 395–7340.

Title: 25 CFR Part 271, Contracts under Indian Self-Determination Act— Subpart H.

Abstract: Any tribe or tribal organization is eligible to contract education programs from the Bureau of Indian Affairs. These rules establish criteria under which a tribe may apply for a new school start or program expansion. The information collected will be in addition to that required under 25 CFR 271.14.

Bureau Form Number: Not applicable. Frequency: Upon initial application. Description of Respondents: Tribes and tribal organizations.

Annual Responses: 2.
Annual Burden Hours: 112 hours.
Bureau Clearance Officer: Cathie
Martin, (202) 343–3577.

Ronal D. Eden,

Acting Deputy to the Assistant Secretary/ Director—Indian Affairs (Indian Education Programs).

[FR Doc. 87-2003 Filed 8-31-87; 8:45 am]
BILLING CODE 4310-02-M

Information Collection Submitted for Review to the Office of Management and Budget for Review Under the Paperwork Reduction Act

The proposal for the collection of information listed below has been submitted to the Office of Management and Budget for approval under the provisions of the Paperwork Reduction Act (44 U.S.C. Chapter 35). Copies of the proposed collection of information and related forms and explanatory material may be obtained by contacting the Bureau's Clearance Officer at the phone number listed below. Comments and suggestions on the requirement should be made within 30 days to the Bureau clearance officer and to the Office of Management and Budget Interior Desk Officer, Washington, DC 20503, telephone (202) 395-7340.

Title: Subchapter E—Education, Adult Education Needs Assessment, 25 CFR Part 46.

Abstract: The Office of Indian Education Programs needs and uses this form as a tool for information gathering and accountability for program integrity while performing its mission of advancing educational opportunities for Native American Indian Adults. Respondents are individuals who seek educational opportunities below the college level.

Bureau Form Number: BIA 62124. Frequency: Annually.

Description of Respondents: Indian/ Alaskan Native students applying for benefits from the Adult Education Program.

Annual Responses: 70.
Annual Burden Hours: 140.
Bureau Clearance Officer: Cathie
Martin, (202) 343–3577.

Mary Widenhouse,

Acting Deputy to the Assistant Secretary/ Director—Indian Affairs (Indian Education Programs).

[FR Doc. 87-20032 Filed 8-31-87; 8:45 am]

Information Collection Submitted for Review to the Office of Management and Budget for Review Under the Paperwork Reduction Act

The proposal for the collection of information listed below has been submitted to the Office of Management and Budget for approval under the provisions of the Paperwork Reduction Act (44 U.S.C. Chapter 35). Copies of the proposed collection of information and related forms and explanatory material may be obtained by contacting the Bureau's Clearance Officer at the phone number listed below. Comments and suggestions on the requirement should be made within 30 days to the Bureau clearance officer and to the Office of Management and Budget Interior Desk Officer, Washington, DC 20503. telephone (202) 395-7340. Title: Subchapter E-Education, Adult Education Annual Report, 25 CFR Part

Abstract: The Office of Indian
Education Programs needs and uses this
form as a tool for information gathering
and accountability for program integrity
while performing its mission of
advancing educational opportunities for
Native American Indian Adults.
Respondents are individuals who seek
educational opportunities below the
college level.

Bureau Form Number: BIA 62123. Frequency: Annually.

Description of Respondents: Indian/ Alaskan Native students applying for benefits from the Adult Education Program.

Annual Responses: 70.
Annual Burden Hours: 140.
Bureau Clearance Officer: Cathie
Martin, (202) 343–3577.

Mary Widenhouse,

Acting Deputy to the Assistant Secretary/ Director—Indian Affairs (Indian Education Programs).

[FR Doc. 87-20033 Filed 8-31-87; 8:45 am] BILLING CODE 4310-02-M

Information Collection Submitted for Review to the Office of Management and Budget for Review Under the Paperwork Reduction Act

The proposal for the collection of information listed below has been submitted to the Office of Management and Budget for approval under the provisions of the Paperwork Reduction Act (44 U.S.C. Chapter 35). Copies of the proposed collection of information and related forms and explanatory material may be obtained by contacting the Bureau's Clearance Officer at the phone number listed below. Comments and suggestions on the requirement should be made within 30 days to the Bureau clearance officer and to the Office of Management and Budget Interior Desk Officer, Washington, DC 20503, telephone (202) 395-7340.

Title: Subchapter E-Education, Adult Education Application Form, 25 CFR Part 46.

Abstract: The Office of Indian Education Programs needs and uses this form as a tool for information gathering and accountability for program integrity while performing its mission of advancing educational opportunities for Native American Indian Adults. Respondents are individuals who seek educational opportunities below the college level.

Bureau Form Number: BIA 6243. Frequency: Annually.

Description of Respondents: Indian/ Alaskan Native students applying for benefits from the Adult Education Program.

Annual Responses: 12,000. Annual Burden Hours: 720. Bureau Clearance Officer: Cathie Martin, (202) 343-3577.

Mary Widenhouse,

Acting Deputy to the Assistant Secretary/ Director-Indian Affairs (Indian Education Programs).

[FR Doc. 87-20034 Filed 8-31-87; 8:45 am] BILLING CODE 4310-02-M

Bureau of Land Management

[NV-930-07-4212-24 (N-33513)]

Airport Lease; Termination of Segregative Effect, Nevada

AGENCY: Bureau of Land Management. Interior.

ACTION: Notice: Termination of Segregative Effect, Nevada.

SUMMARY: This notice terminates the segregative effect of airport lease application, N-33613.

EFFECTIVE DATE: October 1, 1987.

FOR FURTHER INFORMATION CONTACT: Ben Collins, District Manager, Las Vegas District Office, P.O. Box 26569, Las Vegas, NV 89126, [702] 388-6403.

SUPPLEMENTARY INFORMATION: Pursuant to 43 CFR 2091.3-2(b), the Bureau of Land Management hereby terminates the segregative effect as it pertains to the following described lands:

Mount Diablo Meridian, Nevada T. 13 S., R. 71 E.,

Sec. 10, lots 7, 8 and 9.

The airport lease application was filed on June 24, 1981, at which time the lands became segregated from all forms of appropriation. A 20-year lease was subsequently issued to Vincent Silvestri for public airport purposes pursuant to the Act of May 24, 1928 (49 U.S.C. 211 through 214). The facilities were not developed in accordance with section 2(C) of the lease. Furthermore, the Federal Aviation Administration has determined that the site would only support a private use facility. Therefore, Public Airport Lease N-33613 has been cancelled.

At 10:00 a.m., on October 1, 1987, the land will be open to the operation of the public land laws, subject to valid existing rights. All valid applications received prior to or at 10:00 a.m., on October 1, 1987, will be considered as simultaneously filed. All other applications received will be considered in the order of filing.

At 10:00 a.m., on October 1, 1987, the land will also be open to the operation of the mining laws.

Appropriation of lands under the general mining laws prior to the date and time of restoration is unauthorized. Any such attempted appropriation, including attempted adverse possession under 30 U.S.C., 38, shall vest no rights against the United States. Acts required to establish a location and to initiate a right of possession are governed by State law where not in conflict with Federal law. The Bureau of Land Management will not intervene in disputes between rival locators over possessory rights since Congress has provided for such determination in local

The land reimains open to mineral leasing and material sale laws.

Edward F. Spang,

State Director, Nevada.

[FR Doc. 87-20003 Filed 8-31-87; 8:45 am] BILLING CODE 4310-JC-My

[OR-100-84-6310-02: GP-2591

Roseburg District: Advisory Council Meeting

AGENCY: Bureau of Land Management, Interior.

ACTION: Notice of meeting.

SUMMARY: The Bureau of Land Management's Roseburg District Advisory Council will tour the Red Pond area in the North Umpqua Resource Area where they will discuss various land management alternatives.

DATE: September 25, 1987 at 8:15 a.m.

ADDRESS: Bureau of Land Management, 777 NW Garden Valley Blvd., Roseburg. Oregon 97470.

FOR FURTHER INFORMATION CONTACT: Larry Lee, BLM Roseburg District Office, 777 NW Garden Valley Blvd., Roseburg, Oregon 97470. (Telephone (503) 672-4491, Ext. 230.)

SUPPLEMENTARY INFORMATION: The tour is opened to the public and time will be provided for public comment. Members of the public are responsible for their own transportation...

Summary minutes of the meeting will be maintained at the District Office and wil be available for public inspection and reproduction within 30 days following meeting.

M.D. Berg,

District Manager.

Date: August 24, 1987.

[FR Doc. 87-20002 Filed 8-31-87; 8:45 am] BILLING CODE 4310-33-M

[OR080 6310-12 GP7-273]

Salem District; Advisory Council Meeting

ACTION: Notice of Salem District Advisory Council Meeting.

SUMMARY: Notice is hereby given in accordance with section 309 of the Federal Land Policy and Management Act of 1976 that a meeting of the Salem District Advisory Council will be held September 28, 1987, at 1:00 p.m. at the Bureau of Land Management, Salem District Office, 1717 Fabry Road SE., Salem, Oregon.

Agenda for the meeting will include:

- 1. Input into proposed State Director guidance elements in the Bureau of Land Management 1990's Planning for the Public Lands in Western Oregon.
- 2. Program reports (limited to three minutes each) on:
- a. Table Rock Wilderness Management Plan.

 b. Walker Creek Water Supply Project.

c. Yaquina Head Outstanding Natural Area Management Plan.

d. Spotted Owl Management. e. Vegetative Management.

f. Miscellaneous Reports.
The meeting is open to the public.
Anyone wishing to make an oral statement must notify the District Manager at the Salem District Office, 1717 Fabry Road SE, Salem, Oregon 97306 by September 24, 1987. Written comments will also be received for the Council's consideration. Summary minutes will be maintained in the District Office and will be available for public inspection and reproduction during regular business hours within 30

days following the meeting. Dated: August 20, 1987.

Van W. Manning,

District Manager.

[FR Doc. 87-20001 Filed 8-31-87; 8:45 am] BILLING CODE 43:0-33-M

[AZ-020-07-4212-13; A-22792]

Realty Action; Exchange of Public Lands, Maricopa and Cochise Counties, AZ

The following described Federal lands have been determined to be suitable for disposal by exchange pursuant to section 206 of the Federal Land Policy and Management Act of 1976, 43 U.S.C. 1716:

GILA AND SALT RIVER BASE AND MERIDIAN, MARICOPA COUNTY, AZ

Township 5 North, Range 1 East,
Section 27, S½SW¼, NW¼SW¼,
S½NE¼SW¼, NW¼NE¼SW¼;
Section 33, N½NE¼;
Section 34, NW¼, NW¼SW¼,
NW¼NE¼SW¼, NW¼NW¼NE¼,
Containing 450 acres.

In exchange for the above described public lands, the United States will acquire private lands described below from Deputy Partners, an Arizona General Partnership.

GILA AND SALT RIVER MERIDIAN COCHISE COUNTY, AZ

Parcel A

Parcel A (1)

That portion of section 17, Township 24 South, Range 22 East of the Gila and Salt River Base and Meridian, Cochise County, Arizona. Lying northwesterly of the San Pedro River as it ran on February 23, 1982.

Parcel A (2)

The southeast quarter and the east half of the southwest quarter of section 18, Township 24 South, Range 22 East of Gila and Salt River Base and Meridian. Cochise County, Arizona, lying north and west of the San Pedro River, as it ran on February 23, 1982.

Parcel A (3)

All of Lot 3 and any portion of Lot 2 of section 19, Township 24 South, Range 22 East of the Gila and Salt River Base and Meridian, Cochise County, Arizona, lying west of the San Pedro River, as it ran on February 23, 1982.

Parcel B

That portion of the northeast quarter of section 4, Township 24 South, Range 22 East of the Gila and Salt River Base and Meridian, Cochise County, Arizona, more particularly described as follows:

Beginning at the north quarter corner of said section 4:

Thence along the west line of the northeast quarter south 0 degrees 00 minutes 21 seconds west, a distance of 1,321.48 feet;

Thence south 89 degrees 58 minutes 38 seconds east, a distance of 660.00 feet;

Thence north 8 degrees 11 minutes 10 seconds east, a distance of 1,335.35 feet to the north line of said section 4;

Thence along said north line west, a distance of 850.00 feet to the point of beginning.

Except therefrom the north 100.00 feet; and except the west 165.00 feet thereof.

Parcel C

Parcel C (1)

All of the following described property in Township 24 South, Range 22 East of the Gila and Salt River Base and Meridian, Cochise County, Arizona:

Section 3: The west half of the northwest quarter, except the north 100.00 feet.

Section 4: The northeast quarter; the southeast quarter of the northwest quarter; the southwest quarter; the west half of the southeast quarter, except the north 100.00 feet of said northeast quarter.

Section 8: The northeast quarter and the south half:

Section 9: The west half.

Except the following described parcels (A) and (B), from said sections, 3, 4, 8 and 9;

(A) Beginning at the north one-quarter corner of said section 4;

Thence along the west line of the northeast quarter south 0 degrees 00 minutes 21 seconds west, a distance of 1,321.48 feet;

Thence south 89 degrees 58 minutes 38 seconds east, a distance of 660.00 feet;

Thence 8 degrees 11 minutes 10 seconds east, a distance of 1,335.35 feet to the north line of said section 4;

Thence along said north line west, a distance of 850.00 feet to the point of beginning.

Except therefrom the north 100.00 feet as dedicated for road purposes.

(B) Beginning at the north one-quarter corner of said section 4;

Thence south 0 degrees 00 minutes 21 seconds west along the west line of the east one-half of section 4, a distance of 1,321.48 feet.

Thence south 89 degrees 58 minutes 38 seconds east, a distance of 660.00 feet;

Thence south, a distance of 330.00 feet;

Thence south 40 degrees 38 minutes 47 seconds west, a distance of 3,045.97 feet to the southeast corner of the northwest quarter of the southwest quarter of section 4;

Thence south 26 degrees 38 minutes 20 seconds west, a distance of 1,476.64 feet to a point on the south line of section 4, said point being south 89 degrees 54 minutes 30 seconds east, a distance of 660.00 feet from the corner of sections 4, 5, 8 and 9;

Thence south 63 degrees 32 minutes 46 seconds west, a distance of 738.40 feet to a point on the east line of section 8, said point being south 0 degrees 11 minutes 19 seconds west, a distance of 330.00 feet from the corner of sections 4, 5, 8 and 9;

Thence south 0 degrees 11 minutes 19 seconds west along the east line of section 8, a distance of 990.43 feet to the southeast corner of the northeast quarter of the nrotheast quarter of section 8;

Thence south 37 degrees 19 minutes 09 seconds west, a distance of 1,659.60 feet;

Thence south 51 degrees 25 minutes 26 seconds west, a distance of 550.00 feet;

Thence north 74 degrees 15 minutes 25 seconds west, a distance of 1,267.56 feet to the center of section 8;

Thence north 0 degrees 11 minutes 25 seconds east, a distance of 2,641.05 feet to the north quarter corner of section 8;

Thence south 89 degrees 57 minutes 44 seconds east, a distance of 1,325.86 feet to the southwest corner of the southeast quarter of the southeast quarter of section 5;

Thence north 0 degrees 03 minutes 00 seconds west, a distance of 1,320.47 feet to the northwest corner of the southeast quarter of the southeast quarter of section 5;

Thence south 89 degrees 57 minutes 37 seconds east, a distance of 1,325.25 feet to the northeast corner of the southeast quarter of the southeast quarter;

Thence north 0 degrees 04 seconds west, a distance of 1,320.42 feet to the east one-quarter corner of section 5;

Thence south 89 degrees 57 minutes 15 seconds east, a distance of 1,324.79 feet to the southwest corner of the southeast quarter of the northwest quarter of section 4:

Thence north 0 degrees 02 minutes 07 seconds west, a distance of 1,320.96 feet to the northwest corner of the southeast quarter of the northwest quarter of section 4;

Thence south 89 degrees 58 minutes 38 seconds east, a distance of 1,325.74 feet to the northeast corner of the southeast quarter of the northwest quarter of section 4;

Thence north 0 degrees 00 minutes 21 seconds east, a distance of 1,321.48 feet to the point of beginning.

Section 17: The north half of the northeast quarter; the southwest quarter of the northeast quarter; the west half, expecting that portion lying west of the San Pedro River.

Section 18: The southeast quarter; the east half of the southwest quarter, excepting that portion lying north and west of the San Pedro River.

Section 19: Lots 1, 2, and 3, excepting that portion lying west of the San Pedro River. Section 20: Lots 3 and 4.

Parcel C (2)

Parcels 28, 37, 62, 63, 64, 65 and 66, as shown on survey entitled "Amended Plat Palominas Ranches, Unit II," in sections 3, 4, 9, 10, 17 and 20, Township 24 South, Range 22 East of the Gila and Salt River Base and Meridian, Cochise County, Arizona, and recorded in Book 3 of Surveys at pages 19, 19A, 19B and 19C, Cochise County Records.

Parcel D

Parcel D(1)

That portion of the southeast quarter of the southeast quarter of section 29, Township 18 South, Range 21 East of the Gila and Salt River Base and Meridian, Cochise County, Arizona, described as follows:

Beginning 55 rods west of the northeast corner of the southeast quarter of the

southeast quarter;

Thence west 25 rods; Thense south 80 rods;

Thence east 25 rods;

Thence northeasterly to the Point of Beginning.

Parcel D (2)

That portion of the northeast quarter of the southeast quarter of section 29, Township 18 South, Range 21 East of the Gila and Salt River Base and Meridian, Cochise County, Arizona described as follows:

Beginning 45 rods west of the northeast corner of the southeast quarter;

Thence west 35 rods;

Thence south 80 rods;

Thence east 25 rods; Thence northeasterly to the Point of Beginning.

Parcel D (3)

Lot 8 in the southeast quarter of the northeast quarter of section 32, Township 18 South, Range 21 East of the Gila and Salt River Base and Meridian, Cochise County, Arizona.

Parcel D (4)

That portion of Lot 1, or the northeast quarter of the northeast quarter of section 32, Township 18 South, Range 21 East of the Gila and Salt River Base and Meridian, Cochise County, Arizona, described as follows:

Beginning at the southeast corner of the south half of the northeast quarter of the northeast quarter;

Thence north 630.90 feet:

Thence north 72° 06' west, 94.08 feet;

Thence west 1229.99 feet;

Thence south 40 rods;

Thence east 80 rods to the POINT OF BEGINNING.

Parcel D (5)

That portion of the northwest quarter of the northeast quarter of the northeast quarter of section 32, Township 18 South, Range 21 West of the Gila and Salt River Base and Meridian, Cochise County, Arizona, described as follows:

Beginning at the northwest corner of the northeast quarter of the northeast quarter;

Thence south 40 rods; Thence east 25 rods;

Thence north 263.95 feet;

Thence north 72° 06' west, 393.94 feet;

Thence north 17° 54' east, 120 feet;

Thence south 72° 06' east, 355.18 feet;

Thence north 269.95 feet;

Thence west 25 rods to the Point of Beginning;

Except that portion, if any, lying within the following:

A portion of the northeast quarter of the northeast quarter of section 32, Township 18 South, Range 21 East of the Gila and Salt River Base and Meridian, Cochise County, Arizona, more particularly described as follows:

Beginning at the northeast corner, whence the northeast corner of section 32 bears north 73° 26' east, 946.8 feet,

Thence south 63.05 feet to intersect the center line of bridge at north 72° 06' west, 28.98 feet from the center line of West Main Pier:

Thence south 63.05 feet to the southeast corner of this parcel;

Thence north 72° 06' west, 393.94 feet to the southwest corner of this parcel;

Thence north 17° 54' east, 120.0 feet to the northwest corner of this parcel;

Thence south 72° 06' east, 355.18 feet to the Place of Beginning.

Parcel D (6)

Lots 5, 8, 9 and 10, section 33, Township 18 South, Range 21 East of the Gila and Salt River Base and Meridian, Cochise County, Arizona; Except the northern 264.32 feet and a 60° by 60' wellsite.

Parcel D (7)

Lots 1, 2, 7 and 8, section 33, Township 18
South, Range 21 East of the Gila and Salt
River Base and Meridian, Cochise County,
Arizona, according to G.L.O. Survey dated
March 15, 1900 in the records of the Bureau of
Land Management, being the same as Patents
recorded in Book 16 of Deeds of Real Estate,
page 483 and Book 69 of Deeds of Real Estate,
page 157, records of Cochise County, Arizona,
which described said land as the northwest
quarter of the northeast quarter, the northeast
quarter of the northeast quarter, and Lots 1
and 2;

Except the north 825 feet thereof.

Parcel E

Township 23 South, Range 22 East, Section 21, E½NE¼, SW¼NE¼.

Parcel F

Township 23 South, Range 22 East, Section 21, NW1/4.

Comprising 2,675 acres, more or less.

The exchange proposal involves all of the exchange proponent's interest in the surface and mineral estate of the private lands and the surface and mineral estate of the public lands. The exchange is consistent with the Bureau's land use planning objectives.

Lands to be transferred from the United States will be subject to the following reservations, terms and conditions:

1. A right-of-way for ditches and canals constructed by the authority of the United States, Act of August 30, 1890, 26 Stat. 391, 43 U.S.C. 945. 2. Right-of-way AR-024000 to Arizona Public Service Company for electric transmission line purposes.

3. Right-of-way AR-010913 to El Paso Natural Gas Company for natural gas pipeline purposes.

 Right-of-way PHX-086584 to Maricopa County Municipal Water Conservation District.

5. All valid existing rights.

The lands to be acquired by the United States from Deputy Partners shall be subject to certain easements, permits, and other encumbrances detailed in Schedule B of the following TransAmerica Title Insurance Policies: F-830286, F-830287, F-830307, F-830514, F-830515, and F-830516.

In accordance with the regulations of 43 CFR 2201.1(b), publication of this Notice shall segregate the affected public lands from appropriation under the public land laws, including the mining law, and from any subsequent land exchange proposals filed by any proponent other than Deputy Partners or their nominee.

This Notice shall also, inaccordance with 43 CFR 2201.1(b), segregate any and all reserved Federal interest in the offered (non-federal surface) lands from appropriation under the mining laws. This segregation shall terminate two years from the date of this publication or as of the date specified in an opening order published in the Federal Register.

The segregation of the described selected lands shall terminate upon inssuance of a document conveying title to such lands or upon publication in the Federal Register of a notice of termination of the segregation, or the expiration of two years from the date of publication, whichever occurs first.

The appraised value of the offered and selected lands are not equal. A cash equalization payment shall be made by Deputy Partners in accordance with 43 CFR 2201.5(c)(2).

For a period of forty-five (45) days from the date of publication of this notice in the Federal Register, interested parties may submit comments to the Phoenix District Manager, Bureau of Land Management, 2015 West Deer Valley Road, Phoenix, Arizona 85027. Objections will be reviewed by the State Director who may sustain, vacate, or modify this realty action. In the absence of any objections, this realty action will become the final determination of the Department of the Interior.

Paul J. Buff,

Acting District Manager.

Date: August 26, 1987. [FR Doc. 87–19999 Filed 8–31–87; 8:45 am] BILLING CODE 4310–32-M [NM-010-07-4112-01]

District Advisory Council Meeting; Albuquerque District, NM

AGENCY: Bureau of Land Management, Interior.

ACTION: Notice of district advisory council meeting.

SUMMARY: The Bureau of Land Management's Albuquerque District Advisory Council will meet on September 21 and September 22, 1987.

On September 21, the Council will meet at 10:00 a.m. at the Post Office in Lindreth, New Mexico to begin a field tour of BLM roads in the Lindreth and Largo Canyon Areas. The tour will conclude in Farmington, New Mexico at 5:00 p.m.

On September 22, the Council will meet at San Juan College in Farmington in the Sun Dining Hall at 8:30 a.m. Topics to be discussed include possible recommendations on the BLM's road policy and review the Wild Rivers Recreation Area Management Plan.

Members of the public wishing to accompany the Council on the tour must provide their own transportation. Time will be made available for public comments at the September 22 meeting.

The Albuquerque District Advisory
Council is managed in accordance with
the Federal Land Policy and
Management Act of 1979. Minutes of the
meeting will be made available for
review within 30 days following the
meeting. For additional information,
contact Alan Hoffmeister, Public Affairs
Specialist, 435 Montano NE.,
Albuquerque, New Mexico 87107, [505]
766–4504.

Michael Reitz,

Acting District Manager.
[FR Doc. 87-20035 Filed 8-31-87; 8:45 am]
BILLING CODE 4319-FB-M

[UT-920-07-4121-10]

Uinta Southwestern Utah Regional Coal Team Meeting; Utah and Colorado

AGENCY: Bureau of Land Management,

ACTION: Notice of Regional Coal Team Meeting and availability of long range coal market analysis for the Uinta Southwestern Utah Coal Region.

SUMMARY: In accordance with the responsibility outlined in the Federal Coal Management regulations (43 CFR Part 3400), the Regional Coal Team for the Uinta Southwestern Utah Federal Coal Production Region will hold a meeting to review and discuss the status

and need for new Federal Coal Leasing and method of leasing, long range coal market analysis for the Region, draft data adequacy standards, and to discuss other related matters, if any.

The Regional Coal Team is also requesting public comment on subjects to be discussed at the meeting. A copy of the Uinta Southwestern Utah Region Long Range Coal Market Analysis is available from the BLM State Office, Public Room in Utah (Salt Lake City) and Colorado (Denver).

DATE: The Regional Coal Team will meet on October 27, 1987, at 9:00 am.

ADDRESS: The meeting will be held at the Salt Lake Sheraton Hotel, Salon I, 255 South West Temple, Salt Lake City, Utah. Any written comments on the subject to be covered should be provided the BLM State Director, Utah, by October 20, 1987.

FOR FURTHER INFORMATION CONTACT: Max Nielson, Uinta Southwestern Utah Project Manager, Utah State Office, 324

South State Street, Suite 301, Salt Lake City, Utah 84111–2303, telephone [801–524–3004].

SUPPLEMENTARY INFORMATION: The meeting will introduce new members of the recently chartered Regional Coal Team and will discuss recent trends in the region concerning coal leasing and focus on effectively meeting future leasing needs.

Ronald G. Robison,

State Director, Utah.

[FR Doc. 87-19998 Filed 8-31-87; 8:45 am]

National Park Service

National Register of Historic Places; Pending Nominations

Nominations for the following properties being considered for listing in the National Register were received by the National Park Service before August 22, 1987. Pursuant to § 60.13 of 36 CFR Part 60 written comments concerning the significance of these properties under the National Register criteria for evaluation may be forwarded to the National Register, National Park Service, U.S. Department of the Interior, Washington, DC 20243. Written comments should be submitted by September 16, 1987.

Amy Schlagel,

Acting Chief of Registration, National Register.

FLORIDA

Leon County

Tallahassee, Johnson—Carter House, 800 N. Calhoun St.

Pinellas County

Largo, Johnson, Louis, Building, 161 First St., SW.

IDAHO

Adams County

Council, Adams County Courthouse (County Courthouses in Idoho MPS), Michigan St.

Jerome County

Jerome, Jerome County Courthouse (County Courthouses in Idaho MPS), N. Lincoln

Power County

American Falls, Power County Courthouse (County Courthouses in Idaho MPS), Bannock Ave.

Washington County

Weiser, Washington County Courthouse (County Courthouses in Idaho MPS), E. Court St.

IOWA

Dallas County

Earlham vicinity, Wilson, John, House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), S side I-80

Madison County

Earlham vicinity, Early, John and Elizabeth McMurn, House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), 1 mi. S of G31 between P53 & P57

Earlham vicinity, Ford, W. T., House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), 2½ mi. S of Earlham

Earlham vicinity, Henderson, Daniel and Nancy Swaford, House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), 8 mi. S of Earlham on P57

Earlham vicinity, McQuie, Peter and Isabelle McCulloch, Milkhouse (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), SW of Earlham

Earlham vicinity, Seerley, William, Stone Building (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), SE of Earlham

Earlham vicinity, Stone, James Allen, Barn (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), 2½ mi. SE of Earlham

Earlham vicinity, Wilson, Seth and Elizabeth, House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), SE of Earlham

Peru vicinity, Ogburn, William, House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), 11/2 mi. N of E. Peru

Peru, Reed Quarry (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), CR G68, W city limits of Peru

St. Charles vicinity, Holmes, John S. and Elizabeth Beem, Barn (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), CR G50, 1 mi. S.

St. Charles vicinity, Queen, Hogan and Martha A. Runkle, House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), 5 mi. W of St. Charles on CR G50

Winterset vicinity, Armstrong, George and Susan Guiberson, House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), 2½ mi. N of Winterset on G4R

Winterset vicinity, Bevington, C. D. and Eliza Heath, Privy (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), 805 S. Second Ave.

Winterset vicinity, Drake, John and Amanda Bigler, House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), 11 mi. W of Winterset on IA 92

Winterset vicinity, Duff Barn (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), N of Winterset on US 169

Winterset vicinity, Duncan, John E., House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), ½ mi. S of Winterset of P69

Winterset vicinity, Evans, Henry and Elizabeth Adkinson, House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), ½ mi. E of US 169 on CR G50

Winterset vicinity, Macumber, John Andrew and Sara, Ice House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), On G53 1½ mi. E of jct. with P69

Winterset vicinity, McDonald House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), W of Winterset off IA 92 Winterset vicinity, Nichols, William Anzi,

Winterset vicinity, Nichols, William Anzi, House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), E of Winterset of IA 92

Winterset vicinity, Schnellbacher, John and Fredericka, House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), G47, 1½ mi. E of jct. with P53

Winterset vicinity, Seymour Church House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), US 169

Winterset vicinity, Smith, Hiram C., House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), 6 mi. W of Winterset on IA 92

Winterset vicinity, Smith, Hiram C., Milking Shed (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), 6 mi. W of Winterset on IA 92

Winterset, Hornback, Emily, House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), 605 N. First St.

Winterset, Shriver, William R. and Martha Foster, House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), 616 E. Court Ave.

Winterset, Sprague, Brown, and Knowlton Store (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), First & Court Winterset, Vawter, J. G. and Elizabeth S.,

House (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), 223 S. First St.

Winterset, White, Munger and Co. Store (Legacy in Stone: The Settlement Era of Madison County, Iowa TR), 102 W. Court

NEVADA

Carson (Independent City)

Buildings (DeLongchamps, Frederick J., Architecture TR), Carson St.

Clark County

Las Vegas, Railroad Cottage Historic District (Properties Associated with the San Pedro, Los Angeles, and Salt Lake Railroad TR), 601–629 S. Casino Center

NORTH CAROLINA

Cumberland County

Fayetteville, Orange Street School, 500 blk. of Orange St., jct. of Orange & Chance Sts.

NORTHERN MARIANA ISLANDS

Island of Saipan

Chalan Galaide

OHIO

Cuyahoga County

Cleveland, Cleveland Municipal Stadium, Erieview Dr.

Ottawa County

Marblehead, Clemons, Alexander, House, 133 Clemons St.

Stark County

East Canton, Werner Inn, 131 E. Nassau St.

Summit County

Akron, Barder, Byron R., House, 1041 W. Market St.

TEXAS

Brazos County

Bryan, Allen Academy Memorial Hall (Bryan MRA), 1100 blk. Ursuline

Bryan, Allen Block (Bryan MRA), 400-422 N. Main

Bryan, Allen, R.O., House-Allen Academy (Bryan MRA), 1120 Ursuline

Bryan, Armstrong House-Allen Academy (Bryan MRA), 1200 Ursuline

Bryan, Astin, R.Q., House (Bryan MRA), 508 W. Twenty-sixth

Bryan, Blazek, W.J., House (Bryan MRA), 409 W. Thirtieth

Bryan, Bryan Compress and Warehouse (Bryan MRA), 911 N. Bryan

Bryan, Bryan Ice House (Bryan MRA), 107 E.
Martin Luther King

Bryan, CSPS Lodge-Griesser Bakery (Bryan MRA), 304 N. Logan

Bryan, Chance, James O., House (Bryan MRA), 102 S. Parker

Bryan, East Side Historic District (Bryan MRA), Roughly bounded by Houston, Twenty-ninth, Haswell, and E. Thirtieth Sts.

Bryan, Edge, Eugene, House (Bryan MRA), 609 S. Ennis

Bryan, English-Dansby House (Bryan MRA), 204 W. Twenty-eighth

Bryan, English-Poindexter House (Bryan MRA), 206 W. Twenty-eighth

Bryan, First Baptist Church (Bryan MRA), 201 S. Washington

Bryan, First National Bank and Trust Building (Bryan MRA), 120 N. Main

Bryan, First State Bank and Trust Building (Bryan MRA), 100 W. Twenty-fifth Bryan, Higgs, Walter J., House (Bryan MRA),

609 N. Tabor Bryan, House At 1401 Baker (Bryan MRA),

Bryan, House At 1401 Baker (Bryan MRA) 1401 Baker

Bryan, House at 109 N. Sterling (Bryan MRA), 109 N. Sterling

Bryan, House at 407 N. Parker (Bryan MRA), 407 N. Parker

Bryan, House at 600 N. Washington (Bryan MRA), 600 N. Washington

Bryan, House at 603 E. Thirty-first (Bryan MRA), 603 E. Thirty-first Bryan, House at 604 E. Twenty-seventh (Bryan MRA), 604 E. Twenty-seventh Bryan. Humpty Dumpty Store (Bryan MRA), 218 Bryan

Bryan, Jenkins, Edward J., House (Bryan MRA), 607 E. Twenty-seventh

Bryan, Jones, J.M., House (Bryan MRA), 812 S. Ennis

Bryan, Kemp, E.A., House (Bryan MRA), 606 W. Seventeenth

Bryan, McDougal-Jones House (Bryan MRA), 600 E. Twenty-seventh

Bryan, Moore House (Bryan MRA), 500 E. Twenty-fifth

Bryan, Noto House (Bryan MRA), 900 N. Parker

Bryan, Oliver, Dr. William Holt, House (Bryan MRA), 602 W. Twenty-sixth Bryan, Parker Lumber Company Complex

(Bryan MRA), 419 N. Main Bryan, Parker, Milton, House (Bryan MRA),

200 S. Congress
Bryan, Sausley House (Bryan MRA), 700 N.

Washington

Proof. Single Station (Old) (Resear MPA)

Bryan, Sinclair Station, (Old) (Bryan MRA), 507 S. Texas

Bryan, Smith-Barron House (Bryan MRA), 100 S. Congress

Bryan, St. Andrew's Episcopal Church (Bryan MRA), 217 W. Twenty-sixth

Bryan, St. Anthony's Catholic Church (Bryan MRA), 306 S. Parker

Bryan, Stone, Roy C., House (Bryan MRA), 715 E. Thirty-first

Bryan, Zimmerman, Minnie Zulch, House (Bryan MRA), 308 N. Washington

The 15-day commenting period for the following property has been waived in order to assist in the buildings preservation.

NEW YORK

New York County

New York, *United States Courthouse*, 40 Foley Square

[FR Doc. 87-19938 Filed 8-31-87; 8:45 am] BILLING CODE 4310-70-M

Intent To Negotiate Concession Permit; Teton Boating Co., Inc.

Pursuant to the provisions of section 5 of the Act of October 9, 1965 (79 Stat. 969; 16 U.S.C. 20), public notice is hereby given that sixty (60) days after the date of publication of this notice, the Department of the Interior, through the Director of the National Park Service, proposes to negotiate a concession permit with Teton Boating Company. Inc. authorizing it to provide boat transportation and boat rental services for the public at Grand Teton National Park, Wyoming for a period of five (5) years from January 1, 1987, through December 31, 1991.

This permit renewal has been determined to be categorically excluded from the procedural provisions of the National Environmental Policy Act and

no environmental document will be

prepared.

The foregoing concessioner has performed its obligations to the satisfaction of the Secretary under an existing contract which expired by limitation of time on December 31, 1986, and therefore, pursuant to the Act of October 9, 1965, as cited above, is entitled to be given preference in the renewal of the permit and in the negotiation of a new permit as defined in 36 CFR 51.5.

The Secretary will consider and evaluate all proposals received as a result of this notice. Any proposal, including that of the existing concessioner, must be postmarked or hand delivered on or before the sixtieth (60th) day following publication of this notice to be considered and evaluated.

Interested parties should contact the Regional Director, Rocky Mountain Region, P.O. Box 25287, Denver, Colorado 80225, for information as to the requirements of the proposed permit. Jack W. Neckels,

Acting Regional Director, Rocky Mountain Region.

Date July 9, 1987.

[FR Doc. 87-20009 Filed 8-31-87; 8:45 am] BILLING CODE 4310-70-M

[A18 (GUIS-S)]

Gulf Islands National Seashore; Advisory Committee Meeting

July 29, 1987

AGENCY: National Park Service, Gulf Islands National Seashore, Interior. ACTION: Notice of advisory commission meeting.

SUMMARY: Notice is hereby given in accordance with the Federal Advisory Commission Act that a meeting of the Gulf Islands National Seashore Advisory Commission will be held at 2:00 p.m., at the following location and date.

DATE: September 22, 1987.

ADDRESS: William Colmer Visitor Center, Davis Bayou, Ocean Springs, Mississippi.

FOR FURTHER INFORMATION CONTACT: Mr. Jerry Eubanks, Superintendent, Gulf Islands National Seashore, P.O. Box 100, Gulf Breeze, Florida 32561, Telephone: (904) 932–6316.

SUPPLEMENTARY INFORMATION:

The purpose of the Gulf Islands National Seashore Advisory Commission is to consult and advise with the Secretary of the Interior or his designee on matters of planning and development of Gulf Islands National Seashore. The members of the Advisory Commission are as follows:

Mrs. Courtney Blossman, Chairman (Mississippi)

Mr. Gorden D. Allen (Mississippi) Mr. George Byars (Mississippi) Mr. Lloyd Caillavet (Mississippi)

Dr. Ed Cake (Mississippi) Mr. William H. Creel, Sr. (Mississippi)

Mr. Bill Davis (Mississippi)

Mr. Paul Delcambre, Sr. (Mississippi) Ms. Betty S. Goodwin (Mississippi) Mrs. Leewynn Hodges (Mississippi) Mrs. Sara McGehee (Mississippi)

Mr. James E. Walker, Sr. (Mississippi)

Mrs. Lois Anderson (Florida) Mr. Sherman Barnes (Florida)

Mr. J. Earle Bowden (Florida) Mr. Lamar B. Cobb (Florida)

Mr. Paul A. Daniel (Florida) Mrs. Betty Gerritz (Florida)

Mr. Michael Mitchell (Florida) Mrs. Dianne Rittenhouse (Florida)

Mr. Roger Taylor Robinson (Florida) Mr. Walter Francis Spence (Florida)

Mr. Britton Stamps (Florida) Mr. Vince Whibbs (Florida)

The matters to be discussed at this meeting will include:

(1) The status of Park development

(1) The status of Park development plans.

(2) Resource Management and Research Projects update.

(3) Channel maintenance projects. The meeting will be open to the public. However, facilities and space for accommodating members of the public are limited and it is expected that not more than 25 persons will be able to attend. Any member of the public may file with the commission a written statement concerning the matters to be discussed. Written statements may also be submitted to the Superintendent at the address above. Minutes of the meeting will be available at Park Headquarters for public inspection approximately 4 weeks after the meeting.

Robert M. Baker,

Regional Director, Southeast Region.

Date: August 13, 1987.

[FR Doc. 87-20010 Filed 8-31-87; 8:45 am]

BILLING CODE 4310-70-M

INTERSTATE COMMERCE COMMISSION

[No. MC-C-30038]

Petition for Declaratory Order; Extension of Time to File Comments; American Coach Lines, Inc.

AGENCY: Interstate Commerce Commission.

ACTION: Extension of time to file comments.

SUMMARY: By a decision served August 5, 1987, the Commission instituted a proceeding to determine whether incidental charter rights authorize service entirely within the Washington, DC, Metropolitan Area. Notice of the action was published August 6, 1987, in the Federal Register at 52 FR 29317, and in the ICC Register. The due date for comments was set as September 8, 1987. Pursuant to the request of the Washington Metropolitan Area Transit Commission, the time for filing comments has been extended until October 14, 1987.

DATE: Comments may be filed on or before October 14, 1987.

ADDRESSES: Send an original and 10 copies of comments referring to No. MC-C-30038 to: Office of the Secretary, Case Control Branch, Interstate Commerce Commission, Washington, DC 20423.

Send one copy of comments to each of petitioner's representatives:

Leonard A. Jaskiewicz, 1730 M Street, NW., Suite 501, Washington, DC 20036 Lawrence E. Lindeman, 805 King Street, Suite 400, Alexandria, VA 22314

FOR FURTHER INFORMATION CONTACT:

James L. Brown, (202) 275-7898

OF

Andrew L. Lyon, (202) 275-7691

By the Commission, Paul H. Lamboley, Acting Chairman.

Noreta R. McGee,

Secretary.

[FR Doc. 87-20068 Filed 8-31-87; 8:45 am] BILLING CODE 7035-01-M

DEPARTMENT OF LABOR

Office of the Secretary

Office of Administrative Law Judges, Amended Procedures for Internal Handling of Complaints of Misconduct or Disability

AGENCY: Office of Administrative Law Judges, Office of the Secretary, Labor. ACTION: Notice of revision of amended procedures for internal handling of complaints of misconduct or disability.

SUMMARY: Notice is given of revision of the amended procedures for the internal handling of complaints of judicial misconduct or disability on the part of Department of Labor Administrative Law Judges through the establishment of an Advisory Committee. Notice was originally published in the Federal Register on May 22, 1981 (46 FR 28050); and as amended on July 5, 1983 (48 FR 30843).

EFFECTIVE DATE: September 1, 1987.

FOR FURTHER INFORMATION CONTACT: John Vittone, Associate Chief Judge, Office of Administrative Law Judges, 1111 20th Street, NW., Washington, DC 20036, Phone 202–653–5057.

SUPPLEMENTARY INFORMATION: The Amended Procedures for Internal Handling of Complaints of Misconduct or Disability on the part of Department of Labor Administrative Law Judges are revised in order to accommodate the protocol to changes in the number, size and location of our regional or satellite offices. The revision is set forth below.

Subsection 1. of the Procedures section is revised to read as follows:

1. On receipt of one or more written complaints of misconduct or disability of any DOL Administrative Law Judge. or upon his own written charge of misconduct or disability, the Chief Judge shall refer such complaint(s) or charge for limited informal inquiry to an Advisory Committee of three members appointed by him for the particular inquiry from a panel of six judges elected for a term of one year by the DOL Administrative Law Judges other than those who may elect panels in regional offices as hereinafter provided. The judge complained of or charged shall have the right to object to any one or more members of the Committee, in which case such member(s) shall be replaced by one or more judges selected by the Chief Judge from the elected panel.

The Judges in each Regional Office having on its roster ten (10) or more judges in addition to its District Chief Judge (e.g., San Francisco, Cincinnati, and Fort Lauderdale), shall elect a panel of three (3) of its judges from which the Chief Judge shall, in appropriate circumstances and in the manner described above, appoint two (2) judges to serve as a Regional Advisory Committee to function with respect to complaints or charges of misconduct or disability of Administrative Law Judges in such office. A judge elected to any panel (Headquarters or Regional) shall serve thereon for one (1) year or until his/her successor is elected. No District Chief Judge may serve on any panel.

Complaints or charges of misconduct or disability on the part of any Administrative Law Judge employed in a regional or satellite office having less than ten (10) judges (other than the District Chief Judge) on its roster shall be referred, as appropriate, to the (Headquarters) Advisory Committee.

Publication in Final

Notice of proposed rule making has not been published in the Federal

Register as the procedure established herein is a rule of agency practice for which notice and comment is not required. See 5 U.S.C. 553(b)(A).

For the reasons set out in the preamble, the Amended Procedures for Internal Handling of Complaints of Judicial Misconduct are revised as hereinabove set forth.

Signed at Washington, DC on this 19th day of August, 1987.

Nahum Litt.

Chief Judge.

[FR Doc. 87–20071 Filed 8–31–87; 8:45 am] BILLING CODE 4510–20–M

Agency Recordkeeping/Reporting Requirements Under Review by the Office of Management and Budget

Background

The Department of Labor, in carrying out its responsibilities under the Paperwork Reduction Act (44 U.S.C. Chapter 35), considers comments on the reporting and recordkeeping requirements that will affect the public.

List of Recordkeeping/Reporting Requirements Under Review

As necessary, the Department of Labor will publish a list of the Agency recordkeeping/reporting requirements under review by the Office of Management and Budget (OMB) since the last list was published. The list will have all entries grouped into new collections, revisions, extensions, or reinstatements. The Departmental Clearance Office will, upon request, be able to advise members of the public of the nature of the particular submission they are interested in. Each entry may contain the following information:

The Agency of the Department issuing the recordkeeping/reporting requirement.

The title of the recordkeeping/ reporting requirement.

The title of the recordkeeping/ reporting requirement.

The OMB and Agency identification numbers, if applicable.

How often the recordkeeping/ reporting requirement is needed.

Who will be required to or asked to report or keep records.

Whether small businesses or organizations are affected.

An estimate of the total number of hours needed to comply with the recordkeeping/reporting requirement.

The number of forms in the request for approval, if applicable.

An abstract describing the need for and uses of the information collection.

Comments and Questions

Copies of the recordkeeping/reporting requirements may be obtained by calling the Departmental Clearance Officer, Paul E. Larson, telephone (202) 523-6331. Comments and questions about the items on this list should be directed to Mr. Larson, Office of Information Management, U.S. Department of Labor, 200 Constitution Avenue, NW, Room N-1301, Washington, DC 20210. Comments should also be sent to the Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for (BLS/DM/ ESA/ETA/OLMS/MSHA/OSHA/ PWBA/VETS), Office of Management and Budget, Room 3208, Washington, DC 20503 (Telephone (202) 395-6880).

Any member of the public who wants to comment on a recordkeeping/reporting requirement which has been submitted to OMB should advise Mr. Larson of this intent at the earliest possible date.

Extension

Bureau of Labor Statistics

Permanent Mass Layoff and Plant Closing Program

Reports 1–3 and Supplemental Employer Information Report 1220–0090; BLS 428 Quarterly

State or local governments: businesses or other for-profit organizations; Federal agencies or employees; nonprofit institutions.

15,504 responses; 168,055.2 hours; 4 forms

Section 462(e) of the Job Training
Partnership Act states that the
Secretary of Labor develop and
maintain statistical data on
permanent mass layoffs and plant
closings, and publish a report
annually. These data will be used to
study the causes and effects of worker
dislocations.

Employment and Training Administration

Interstate Arrangement for Combining Employment and Wages 1205–0029; ETA 586

Quarterly
State or local governments
53 respondents; 636 hours; 1 form
3304(a)(9)(B) of the I.R.C. of 1954
requires States to participate in
arrangement, as prescribed by the
Secretary of Labor, which provide for
the payment of UI benefits on the
basis of combining employment and

the payment of UI benefits on the basis of combining employment and wages earned in two or more states. This report is needed to measure the scope and monitor the operation of this program.

Occupational Safety and Health Administration

Ethylene Oxide 1218–0108

Recordkeeping; on occasion Businesses or other for-profit; Federal agencies or employees; Small businesses or organizations

6453 respondents; 215,036 burden hours; 0 forms

This regulation requires employers to establish and maintain accurate records of exposure monitoring and medical surveillance for employees exposed to ethylene oxide (EtO). These records will be used by employers, employees, physicians and the Government to ensure that workplace exposure to EtO does not adversely affect the health of employees.

Signed at Washington, DC this 27th day of August, 1987.

Paul E. Larson,

Departmental Clearance Officer. [FR Doc. 87-20070 Filed 8-31-87; 8:45 am] BILLING CODE 4510-28-M

Employment and Training Administration

Investigations Regarding Certifications of Eligibility To Apply for Worker Adjustment Assistance; Al Tech Specialty Steel Corp., et al.

Petitions have been filed with the Secretary of Labor under section 221(a) of the Trade Act of 1974 ("the Act") and are identified in the Appendix to this notice. Upon receipt of these petitions, the Director of the Office of Trade Adjustment Assistance, Employment and Training Administration, has instituted investigations pursuant to section 221(a) of the Act.

The purpose of each of the investigations is to determine whether the workers are eligible to apply for adjustment assistance under Title II, Chapter 2, of the Act. The investigations will further relate, as appropriate, to the determination of the date on which total or partial separations began or threatened to begin and the subdivision of the firm involved.

The petitioners or any other persons showing a substantial interest in the subject matter of the investigations may request a public hearing, provided such request is filed in writing with the Director, Office of Trade Adjustment Assistance, at the address shown below, not later than September 11, 1987.

Interested persons are invited to submit written comments regarding the subject matter of the investigations to the Director, Office of Trade Adjustment Assistance, at the address shown below, not later than September 11, 1987.

The petitions filed in this case are available for inspection at the Office of the Director, Office of Trade Adjustment Assistance, Employment and Training Administration, U.S. Department of Labor, 601 D Street NW., Washington, DC 20213.

Signed at Washington, DC, this 24th day of August 1987.

Glenn M. Zech,

Acting Director, Office of Trade Adjustment Assistance.

APPENDIX

Petitioner: Union/workers/firm	Location	Date received	Date of petition	Petition	Articles produced
Anchor Hocking/Newell Co. (AFGWU) Dee Cee Apparel (Workers) Grace Shoe Mfg. Co. Inc. (Workers) Holley Auto DivColt Industries (Workers) Imperial Knife Div. (Company) Nicor Drilling Co. (Workers) Sifco Industries (Company) Switches, Inc. (Workers)	Perrysburg, OH Lancaster, OH Hohenwald, TN Somersworth, NH Paris, TN Providence, BI Lafayette, LA Byesville, OH Leiters Ford, IN Midland, TX Pittsburgh, PA	8/24/87 8/24/87 8/24/87 8/24/87 8/24/87 8/24/87 8/24/87 8/24/87 8/24/87 8/24/87 8/24/87	7/29/87 8/17/87 8/11/87 8/6/87 8/10/87 8/12/87 8/13/87 8/10/87 8/14/87 8/12/87 8/11/87	20, 028 20, 029 20, 030 20, 031 20, 032 20, 033 20, 034 20, 035	Tableware. Pants. Shoes. Carburetors. Tableware. Oil and gas. Yokes and gears.

[FR Doc. 87-20072 Filed 8-31-87; 8:45 am] BILLING CODE 4510-30-M

[TA-W-19,671]

Affirmed Determination Regarding Application for Reconsideration; Hobart Corp., Dayton, OH

By an application dated July 27, 1987, the International Union of Electronic and Electrical Workers (IUE) requested administrative reconsideration of the Department of Labor's Notice of Negative Determination Regarding Eligibility to Apply for Worker Adjustment Assistance on behalf of former workers of Hobart Corporation, Dayton, Ohio. The determination was published in the Federal Register on July 21, 1987 (52 FR 27479).

The union claims that Hobart Corporation is transferring the production of food weighing scales from Dayton to Taiwan. According to the union, the transferred production will be sold to Hobart's domestic customers.

Conclusion

After careful review of the application, I conclude that the claim is of sufficient weight to justify reconsideration of the Department of Labor's prior decision. The application is, therefore, granted.

Signed at Washington, DC, this 21st day of August, 1987.

Harold A. Bratt,

Deputy Director, Office of Program Management, UIS.

[TA-W-19.629]

Dismissal of Application for Reconsideration; J.E. Carter Energy & Development Corp., Houston, TX

Pursuant to 29 CFR 90.18 an

application for administrative reconsideration was filed with the Director of the Office of Trade Adjustment Assistance for workers at J.E. Carter Energy & Development Corporation, Houston, Texas. The review indicated that the application contained no new substantial information which would bear importantly on the Department's determination. Therefore, dismissal of the application was issued.

TA-W-19,629; J.E. Carter Energy & Development Corp., Houston, TX (August 25, 1987)

Signed at Washington, DC this 25th day of August 1987.

Glenn M. Zech.

Acting Director, Office of Trade Adjustment Assistance.

[FR Doc. 87-20014 Filed 8-31-87; 8:45 am]
BILLING CODE 4510-30-M

[TA-W-19, 961]

Termination of Investigation; Omnisport Inc. Woonsocket, RI

Pursuant to section 221 of the Trade
Act of 1974, an investigation was
initiated on July 27, 1987 in response to a
worker petition received on July 27, 1987
which was filed by the Amalgamated
Clothing and Textile Workers Union on
behalf of workers producing nylon and
wool athletic and award jackets at
Omnisport Incorporated, Woonsocket,
Rhode Island.

The petitioner, Mr. Steve Stycos, union representative for workers at Omnisport Incorporated, Woonsocket, Rhode Island has requested that the petition be withdrawn by letter, received August 12, 1987. Consequently, further investigation in this case would serve no purpose; and the investigation has been terminated.

Signed at Washington, DC, this 17th day of August 1987.

Marvin M. Fooks,

Director, Office of Trade Adjustment Assistance.

[FR Doc. 87-20075 Filed 8-31-87; am]

Determinations Regarding Eligibility to Apply for Worker Adjustment Assistance; Butler Livestock Systems, et al

In accordance with section 223 of the Trade Act of 1974 (19 U.S.C. 2273) the Department of Labor herein presents summaries of determinations regarding eligibility to apply for adjustment assistance issued during the period August 17, 1987—August 21, 1987

In order for an affirmative determination to be made and a certification of eligibility to apply for adjustment assistance to be issued, each of the group eligibility requirements of section 222 of the Act must be met.

 That a significant number or proportion of the workers in the workers' firm, or an appropriate subdivision thereof, have become totally or partially separated,

(2) That sales or production, or both, of the firm or subdivision have decreased absolutely, and

(3) That increases of imports of articles like or directly competitive with articles produced by the firm or appropriate subdivision have contributed importantly to the separations, or threat thereof, and to the absolute decline in sales or production.

Negative Determinations

In each of the following cases the investigation revealed that criterion [3]

has not been met. A survey of customers indicated that increased imports did not contribute importantly to worker separations at the firm.

TA-W-19.842; Butler Livestock Systems, A Division of Butler Manufacturing Co., Fort Atkinson, WI

TA-W-19,832; Midwest Carbide Corp., Keokuk, IA

In the following cases, the investigation revealed that criterion (3) has not been met for the reasons specified.

TA-W-19,950; Federal Ore and Chemical, Inc., (Formerly Federal Bentonite Div. of Aurora Industries), Culver, MN

U.S. imports of bentonite are negligible.

TA-W-20,005; Double E. Well Service, St. Louis, MO

The workers' firm does not produce an article as required for certification under section 222 of the Trade Act of 1974.

TA-W-19,888; Central Oilfied Supply Co., Wooster, OH

U.S. imports of pipe and tubing declined absolutely in 1986 compared to 1985 and in the January through March period of 1987 compared to the corresponding 1986 period.

TA-W-19.911; American Acceptance Corp., Norristown, PA

The workers' firm does not produce an article as required for certification under section 222 of the Trade Act of 1974.

TA-W-19,886; Arco Oil and Gas Co., Div. of Arco Healdton Suboffice, Healdton, OK

Increased imports did not contribute importantly to workers separations at the firm.

TA-W-19,962; Port Clyde Foods, Inc., Maine Container Div., Rockland, ME

The investigation revealed that criterion (2) has not been met. Sales or production did not decline during the relevant period as required for certifications.

TA-W-20,000; Texas Gas Corp., Corpus Christi

The workers' firm does not produce an article as required for certification under section 222 of the Trade Act of 1974.

TA-W-19,920; Simpson Timber Co., Columbia Door Div., Vancouver, WA

U.S. imports of flush type wood doors declined absolutely and relative to domestic shipments in 1986 compared to 1985 and declined absolutely in the first quarter of 1987 to the first quarter 1986.

TA-W-19,915; Eureka Pipe Line Co., Parkersburg, WV

The workers' firm does not produce an article as required for certification under section 222 of the Trade Act of 1974.

TA-W-20,024; San Antonio Data Processing, Inc., San Antonio, TX

The workers' firm does not produce an article as required for certification under section 222 of the Trade Act of 1974.

Affirmative Determinations

TA-W-19,869; W.E. Stephens, Watertown, TN

A certification was issued covering all workers of the firm separated on or after June 22, 1986.

TA-W-19,851; Gent J. Manufacturing, Inc., Plymouth, PA

A certification was issued covering all workers of the firm separated on or after June 24, 1986.

TA-W-19,817; Par Microsystems Corp., New Hartford, NY

A certification was issued covering all workers of the firm separated on or after June 4, 1986.

TA-W-19,960; Mobil Exploration and Producing, Denver Division, Denver, CO

A certification was issued covering all workers of the firm separated on or after June 25, 1986.

TA-W-19,819; Samco Sportswear, Crosby, MN

A certification was issued covering all workers of the firm separated on or after June 1, 1986 and before June 15, 1987.

TA-W-19,823; Brookevale Manufacturing Co., Belle Vernon, PA

A certification was issued covering all workers of the firm separated on or after March 1, 1987.

TA-W-19,848; E.F. Johnson Co., Waseca, WI

A certification was issued covering all workers of the firm separated on or after June 19, 1986.

TA-W-19,853; Holly Dress Co., Nanticoke, PA

A certification was issued covering all workers of the firm separated on or after June 22, 1986.

TA-W-19.893; FPCO Oil and Gas Co., Houston, TX

A certification was issued covering all workers of the firm separated on or after July 1, 1986. TA-W-19,893A; FPCO Oil and Gas Co., Bakersfield, CA

A certification was issued covering all workers of the firm separated on or after July 1, 1986.

TA-W-19,893B; FPCO Oil and Gas Co., Denver, CO

A certification was issued covering all workers of the firm separated on or after July 1, 1986.

I hereby certify that the aforementioned determinations were issued during the period August 17, 1987-August 21, 1987. Copies of these determinations are available for inspection in Room 6434, U.S. Department of Labor, 601 D Street, NW., Washington, DC 20213 during normal business hours or will be mailed to persons who write to the above address.

Dated: August 25, 1987.

Glenn M. Zech,

Acting Director Office of Trade Adjustment Assistance.

[FR Doc. 87-20076 Filed 8-31-87; 8:45 am] BILLING CODE 4510-30-M

Job Training Partnership Act; Debt Collection; Guidance to States

AGENCY: Employment and Training Administration, Labor.

ACTION: Notice of method of repayment to recover misexpenditures.

SUMMARY: The Employment and Training Administration of the Department of Labor has issued instructions to the States on the recovery of misexpenditures under the Job Training Partnership Act. The notice is given to interested persons in order that they may familiarize themselves with the types of misexpenditures requiring repayment in non-Federal funds and the disposition of recoveries from other less serious misexpenditures. This guidance was provided in a Training and Employment Guidance Letter issued August 11, 1987 and published at the end of this document. EFFECTIVE DATE: August 11, 1987.

FOR FURTHER INFORMATION CONTACT:
Ms. Linda D. Kontnier, Chief, or James
Mac Donald, Division of Debt
Management, Office of Grants and
Contracts Management, Employment
and Training Administration,
Department of Labor, Room N4671, 200
Constitution Avenue, NW., Washington,
DC 20210. Telephone: [202] 535-0704.

SUPPLEMENTARY INFORMATION: On May 12, 1987, the Employment and Training Administration of the Department of Labor published in the Federal Register a notice of a proposed Training and Employment Guidance Letter on the disposition of misexpended funds recovered under the Job Training Partnership Act. Interested parties were invited to submit comments until June 11, 1987. This notice summarizes the comments received and announces the issuance of the Training and Employment Guidance Letter.

Discussion of Comments

A total of eight replies were received. They contained numerous comments in response to the May 12, 1987, Federal Register notice of the proposed guidance letter. In their letters, recipients expressed concerns about the policy and the impact of its implementation.

The comments have been grouped into six categories for purposes of analysis and resolution:

(1) A number of recipients requested additional guidance definition to enable them to identify more precisely those

instances in which it is necessary to remit misexpenditures to the

Department.

At this early stage of the ITPA program, it is not possible to identify every circumstance for which recovered misexpenditures must be returned to the Department. Recipients are referred to sections 164(a) to 164(e)(1) of the Act, the implementing regulations at 20 CFR 629.44, TEGL No. 6-84, and OMB Circular A-128. These guidance materials should be read within the context of DOL's objectives as stated in the background section. First, DOL is committed to vigorous corrective action in instances of serious violations or illegal acts. At the same time, DOL is equally committed to a conservation of funds for program use. While there may appear to be a dichotomy between the two objectives, the long term prospects of achieving the second objective is enhanced by the enforcement of sanctions in those instances in which misexpenditures are due to serious violations or illegal acts.

It will be the responsibility of recipients to exercise judgment in determining if the misexpenditure in question should be categorized as a serious violation or illegal act. While the nature of some cases may be obvious, others may require the recipient's best judgment based on the available facts. Such a judgment should reflect the interests of responsible public policy that is concerned with the integrity of the recipient's employment and training delivery system as a whole. While the practical effect of a repayment on the integrity of the delivery system may be difficult to measure quantitatively, experience has shown that it has both an immediate and longterm effect.

There are no changes in the TEGL as a result of these comments.

(2) Other commenters complained that the net result of the policy will be to reduce the resources available to provide job training services. Relatedly, one commenter argued that the policy acts as a de facto reduction of the JTPA grant which should be effectuated (only) through legislative action. Section 164 of the Act directs the repayment of misexpenditures so that no further legislative action is necessary.

When non-Federal funds are remitted to the Department in settlement of a debt, there is no reduction of JTPA funds and therefore no reduction of the funds available to provide job training services. Only if JTPA funds are remitted to the Department or if the Secretary effects an allocations offset is there a reduction of the resources available to provide job training services.

There are no changes in the TEGL as a result of these comments.

(3) Some commenters suggested that the three year limitation for reprogramming be eliminated or extended.

Section 161(b) of the Act states that, "Funds obligated for any program year may be expended by each recipient during that program year and the two succeeding program years * * *."

This is a standard limitation that the Congress imposes on appropriations in order to maintain control of public funds. Short of amending the legislation, there is no possibility of changing this limitation.

There are no changes in the TEGL as a result of these comments.

(4) A few commenters asked the question whether there will be any provision for repayment of misexpenditures by a method other than non-Federal funds.

The proposed TEGL directs recipients to collect and repay only those amounts associated with misexpenditures which relate to the Department's first objective of vigorous corrective action: serious violations or illegal acts. There are no provisions for repayment of these misexpenditures by a method other than non-Federal funds.

With respect to misexpenditures which are not in the category of "serious violations or illegal acts," the proposed TEGL addresses only those situations in which the subrecipient actually remits non-Federal funds to the recipient. In those instances, the recipient is to reprogram the funds, subject to the 3 year expenditure limitation.

A phrase has been inserted in paragraph seven to make clear that, in

instances where the liability does not arise under one of the serious categories, the TEGL addresses only those situations in which the recipient has recovered non-Federal funds.

(5) One commenter complained that the proposed policy requiring repayment of non-Federal funds was more restrictive than the law and the regulations which the commenter said "permit the use of other Federal funds to repay debts."

In reference to repayment, section 164(e)(1) uses the phrase, "from funds other than funds received under this Act." Such a phrase does not make the affirmative statement that other Federal funds can be used to repay a JTPA misexpenditure. It is the policy of ETA not to accept other Federal funds in payment of debts. To do otherwise would be to encourage, or at least condone, a type of offset that would defeat the purposes of other Federal programs.

There are no changes in the TEGL as a result of this comment.

(6) One commenter asked if there was a relationship between the consideration of the four waiver factors at 164(e)(2)(A-D) of the Act and the determination that a misexpenditure is in the category of "serious violations or illegal acts." There is no relationship between the two. In the process of responding to a request for waiver of the recipient's liability for a subrecipient misexpenditure, the Secretary reviews the four factors in order to determine if the recipient has demonstrated substantial compliance with the factors.

In determining whether a misexpenditure is in the category of "serious violation or illegal act," a recipient reviews the relevant information available, using as background guidance the references in the TEGL, including sections 164(a) to 164(e)(1) of the Act, but not section 164(e)(2) which contains the four factors that the Secretary uses in his determination.

There are no changes in the TEGL as a result of this comment.

Signed at Washington, DC, this 25th day of August 1987.

Roger D. Semerad,

Assistant Secretary of Labor.

Date: August 11, 1987.

Training and Employment Guidance Letter No. 2-87]

From: Roger D. Semerad, Assistant Secretary of Labor.

Subject: Debt Collection Under the Job Training Partnership Act (JTPA).

1. Purpose

To provide States with instructions on the method of repayment to recover misexpenditures under the Job Training Partnership Act.

2. References

Debt Collection Act of 1982 [31 U.S.C. 3701 et seq.); Federal Claims Collection Standards (4 CFR Chapter II): Training and Employment Guidance Letter (TEGL) No. 6-84; OMB Circular A-128; Training and Employment Information Notice (TEIN) No. 7-86.

3. Background

Training and Employment Information Notice (TEIN) No. 7-86 was issued on August 14, 1986, to address a very narrow concern: Insuring consistency in the Employment and Training Administration's (ETA) response to States' inquiries on how to handle misspent ITPA funds recovered from sub-recipients. Since the issuance of TEIN No. 7-86, further consideration has been given to two additional objectives: (1) The reinforcement of the Department of Labor's commitment to vigorous corrective action in relation to instances of serious violations or illegal acts, and (2) the conserving of funds for program

4. Method of Repayment

Section 164(e)(1) of the Job Training Partnership Act requires repayment from non-JTPA funds where the misexpenditure was due to "willful disregard of the requirements of the Act, gross negligence, or failure to observe accepted standards of administration."

For those misexpenditures where the

liability arises from:

(1) "Willful disregard," "gross negligence," or "failure to observe accepted standards of administration,"

under section 164(e)(1);
(2) "Incidents of fraud, malfeasance, misapplication of funds" or other serious violations as defined in TEGL No. 6-84;

(3) "Illegal acts or irregularities" which are required to be reported in accordance with paragraphs 11.b.(4) and 12. of OMB Circular A-128,

it is ETA policy that non-Federal funds must be remitted to the Department of Labor and these amounts will not be available for reprogramming; such funds will revert to the U.S. Treasury. This policy applies whether the misexpenditure occurs at the recipient or at any sub-recipient level. These funds are to be remitted to:

U.S. Department of Labor, **Employment and Training**

Administration, Office of Financial and Administrative Management, Room N 4671, 200 Constitution Avenue, NW., Washington, DC 20210.

The letter of transmittal should provide the following identifying information:

Governor/Secretary Agreement Number (Grant No.), program title, program year to which the repayment applies, sub-recipient name, subrecipient grant or contract number.

In those instances where the liability does not arise under the above categories and there has been a repayment of non-Federal funds to the recipient, the funds shall be reprogrammed into the same JTPA program and title, provided this reprogramming takes place during the program year the funds were obligated by DOL, or the two succeeding program years.

Any funds remitted to ETA because the three year availability period has expired, must be remitted to the Department of Labor at the above address. Such funds will not be available to the State for subsequent drawdown or expenditure.

In those cases where non-Federal funds are reprogrammed, documentation relating to the repayment of the liability and the reprogramming of the funds should be maintained and available for review during the compliance review process.

5. Effective Date

This Training and Employment Guidance Letter shall be effective as of the date of issuance.

6. Rescission.

Training and Employment Information Notice No. 7-86 is cancelled as of the above effective date of this issuance.

7. Promulgation

The guidance contained in this letter will be published in the Federal Register. Copies of the TEGL are being sent to State ITPA liaisons.

8. Inquiries

Questions concerning this guidance letter should be directed to Linda Kontnier or James MacDonald on (202) 535-0704.

Expiration date: August 31, 1988.

[FR Doc. 87-20077 Filed 8-31-87; 8:45 am]

BILLING CODE 4510-30-M

NUCLEAR REGULATORY COMMISSION

[Docket No. 50-293]

Environmental Assessment and Finding of No Significant Impact; Boston Edison Co.

The U.S. Nuclear Regulatory
Commission (the Commission) is
considering issuance of exemptions from
certain requirements of 10 CFR Part 50,
Appendix R, to the Boston Edison
Company (BECo/licensee) for the
Pilgrim Nuclear Power Station located at
the licensee's site in Plymouth County,
Massachusetts.

Environmental Assessment

Identification of the Proposed Action

The proposed action would grant exemptions from certain specific requirements of Appendix R of 10 CFR Part 50. Specifically exemptions were requested from section III. J to the extent that, "Emergency lighting units with at least an 8-hour battery power supply shall be provided in all areas needed for operation of safety shutdown equipment and in access and egress routes thereto."

The exemption is responsive to the licensees request dated April 1, 1987.

The Need for the Proposed Action

The proposed exemption is needed because the features described in the licensee's request regarding the existing yard security lighting are the most practical method for meeting the fire protection emergency lighting requirements of Appendix R. Literal compliance would not significantly enhance the fire protection capability.

Environmental Impacts of the Proposed Action

The proposed exemption will provide a degree of fire protection such that there is no increase in the risk of fires at the facility. Consequently, the probability of fires has not been increased and the post-fire radiological releases will not be greater than previously determined nor does the proposed exemption otherwise affect radiological plant effluents. Therefore, the Commission concludes that there are no significant radiological environmental impacts associated with its proposed exemption.

The proposed exemptions do not affect nonradiological plant effluents and have no other environmental impact. Therefore, the Commission concludes that there are on significant nonradiological environmental impacts

associated with the proposed exemption.

Alternative Use of Resources

This action involves no use of resources not previously considered in the Final Environmental Statement for the Pilgrim Nuclear Power Station.

Agencies and Persons Consulted

The NRC staff reviewed the licensee's request and did not consult other agencies or persons.

Findings of No Significant Impact

The Commission has determined not to prepare an environmental impact statement for the proposed exemption.

Based upon the foregoing environmental assessment, the staff concludes that the proposed action will not have a significant effect on the quality of the human environment.

For further details with respect to this proposed action, see the licensee's letter dated April 1, 1987. This letter is available for public inspection at the Commission's Public Document Room, 1717 H Street, NW., Washington, DC., and at the Plymouth Public Library, 11 North Street, Plymouth County, Massachusetts, 02360.

Dated at Bethesda, Maryland this 26th day of August, 1987.

For the Nuclear Regulatory Commission. Richard H. Wessman,

Acting Director. Project Directorate I-3, Division of Reactor Projects I/II. [FR Doc. 87-20062 Filed 8-31-87; 8:45 am] BILLING CODE 7590-01-M

[Docket No. 50-255]

Environmental Assessment and Finding of No Significant Impact; Consumers Power Co.

The U.S. Nuclear Regulatory
Commission (the Commission) is
considering issuance of an exemption
from the schedular requirements of
Appendix J to 10 CFR Part 50 to
Consumers Power Company (the
licensee), for the Palisades Plant,
located in Van Buren County, Michigan.

Environmental Assessment

Identification of Proposed Action

The exemption would provide an alternative to the requirement of 10 CFR Part 50, Appendix J, section III.A.6(b), to perform at each plant shutdown for refueling or approximately every 13 months, whichever occurs first, a Type A Containment Integrated Leak Rate Test until two consecutive tests meet the acceptance criteria, if two consecutive Type A tests performed in accordance with the normal schedule fail to meet

the criteria. At Palisades, the containment has failed the last three Type A tests because of excess leakage penalties applied from the Type B and C tests for locally testable penetrations. The alternative that the licensee has proposed for the accelerated Type A testing is an accelerated Type B and C testing program, along with aggressive maintenance of these penetrations based on trending information and engineering evaluations of any problems encountered, since the previous Type A failures have always been attributable to the penetrations associated with the local Type B and C tests.

The Need for the Proposed Action

The proposed exemption is needed because the licensee's proposed "Local Leak Rate Testing—Corrective Action Plan" described in the licensee's request is the most practical method of meeting the intent of Appendix J and literal compliance would not serve the underlying purpose of Appendix J and is not necessary to achieve the underlying purpose of Appendix J.

Environmental Impacts of Proposed Action

The proposed exemption would provide a degree of assurance of containment integrity equivalent to that required by Appendix J. Consequently, the probability of accidents has not been increased by this administrative change and the post-accident radiological releases would not be greater than previously determined. Neither does the proposed exemption otherwise affect radiological plant effluents. Therefore, the Commission concludes that there are no significant radiological environmental impacts associated with this proposed exemption.

With regard to potential nonradiological impacts, the proposed exemption involves a change in surveillance or inspection requirements. It does not affect non-radiological plant effluents and has no other environmental impact. Therefore, the Commission concludes that there are no significant non-radiological environmental impacts associated with the proposed exemption.

Alternatives to the Proposed Action

Since the Commission has concluded that the environmental effects of the proposed action are negligible, any alternatives with equal or greater environmental impacts need not be evaluated.

The principal alternative would be to deny the requested exemption. This

would not reduce the environmental impacts attributable to this facility and would result in a larger expenditure of licensee resources to comply with the Commission's regulations.

Alternative Use of Resources

This action involves no use of resources not previously considered in the Final Environmental Statement related to operation of the Palisades Plant.

Agencies and Persons Consulted

The Commission's staff reviewed the licensee's request and did not consult other agencies or persons.

Finding of No Significant Impact

The Commission has determined not to prepare an environmental impact statement for the proposed exemption.

Based upon the foregoing environmental assessment, we conclude that the proposed action will not have a significant effect on the quality of the human environment.

For further details with respect to this action, see the application for exemption dated August 22, 1986, which is available for public inspection at the Commission's Public Document Room, 1717 H Street, NW., Washington, DC, and at the Van Zoeren Library, Hope College, Holland, Michigan 49423.

Dated at Bethesda, Maryland, this 25th day of August 1987.

For the Nuclear Regulatory Commission. Albert W. De Agazio,

Acting Director, Project Directorate III-1, Division of Reactor Projects, III, IV, V and Special Projects.

[FR Doc. 87-20061 Filed 8-31-87; 8:45 am]

Draft Regulatory Guide; Issuance, Availability

The Nuclear Regulatory Commission has issued for public comment a draft of a proposed revision to a guide in its Regulatory Guide Series together with a draft of the associated value/impact statement. This series has been developed to describe and make available to the public such information as methods acceptable to the NRC staff for implementing specific parts of the Commission's regulations, techniques used by the staff in evaluating specific problems or postulated accidents, and data needed by the staff in its review of applications for permits and licenses.

The draft, temporarily identified by its task number, EE 108–5 (which should be mentioned in all correspondence concerning this draft guide), is proposed Revision 2 to Regulatory Guide 1.100, "Seismic Qualification of Electric and

Mechanical Equipment for Nuclear Power Plants." This guide is being revised to provide guidance on a method acceptable to the NRC staff for complying with NRC's regulations with respect to seismic qualification of mechanical as well as electric equipment. This guide endorses, with certain exceptions, the revised IEEE Std 344–1987, "Recommend Practice for Seismic Qualification of Class 1E Equipment for Nuclear Power Generating Stations."

This draft guide and the associated value/impact statement are being issued to involve the public in the early stages of the development of a regulatory position in this area. They have not received complete staff review and do not represent an official NRC staff position.

Public comments are being solicited on both the guide (including any implementation schedule) and the value/impact statement. Comments on the draft value/impact statement should be accomplished by supporting data. Written comments may be submitted to the Rules and Procedures Branch, Division of Rules and Records, Office of Administration and Resources Management, U.S. Nuclear Regulatory Commission, Washington, DC 20555. Comments may also be delivered to Room 4000, Maryland National Bank Building, 7735 Old Georgetown Road, Bethesda, Maryland from 8:15 to 5:00 p.m. Copies of comments received may be examined at the NRC Public Document Room, 1717 H Street NW., Washington, DC. Comments will be most helpful if received by October 30, 1987.

Although a time limit is given for comments on these drafts, comments and suggestions in connection with (1) items for inclusion in guides currently being developed or (2) improvements in all published guides are encouraged at any time.

Regulatory guides are available for inspection at the Commission's Public Document Room, 1717 H Street NW., Washington, DC. Requests for single copies of draft guides (which may be reproduced) or for placement on an automatic distribution list for single copies of future draft guides in specific divisions should be made in writing to the U.S. Nuclear Regulatory Commission, Washington, DC 20555, Attention: Director, Division of Information Support Services. Telephone requests cannot be accommodated. Regulatory guides are not copyrighted, and Commission approval is not required to reproduce them.

(5 U.S.C. 552(a))

Dated at Rockville, Maryland, this 26th day of August 1987.

For the Nuclear Regulatory Commission. Guy A. Arlotto,

Director, Division of Engineering, Office of Nuclear Regulatory Research.

[FR Doc. 87-20060 Filed 8-31-87; 8:45 am]

Reactor Risk Reference Document (NUREG-1150); Meeting of the Peer Review Committee

AGENCY: Nuclear Regulatory Commission.

ACTION: Notice of meeting.

SUMMARY: The draft Reactor Risk Reference Document (NUREG-1150) is currently undergoing a detailed peer review by a fourteen member committee chaired by Dr. William E. Kastenberg of the University of California, Los Angeles. Administrative and technical support is being provided by the Lawrence Livermore National Laboratory (LLNL), funded by the Nuclear Regulatory Commission (NRC). The peer review committee met during June 24-25, 1987 at the LLNL and July 15-17, 1987 at Albuquerque, NM. Three additional meetings are scheduled before the end of the year.

DATES AND TIMES: The third meeting will be held during September 15–17, 1987, from 8:30 am to 5:00 pm on September 15 and 16, and from 8:30 am to 12 noon on September 17.

The fourth and the fifth meetings are scheduled to be held November 9-11 and December 10-11, 1987, respectively.

ADDRESS: The September meeting will be at the Holiday Inn, 10740 Wilshire Blvd., Los Angeles, CA 90024.

The location of the November and December meetings will be announced at a later date.

FOR FURTHER INFORMATION CONTACT: Dr. Pradyot K. Niyogi, Division of Reactor Accident Analysis, Office of Nuclear Regulatory Research, Washington, DC 20555, (301) 443–7611.

SUPPLEMENTARY INFORMATION: Active participation in the meeting will be limited to the members of the committee, but the meeting will be open to the public to attend as observers. Members of the public may submit written comments on topics related to the meeting discussion. Limited verbal comment by the public will be permitted during the meeting at specified times. Prospective attendees should notify Dr. Sergio Guarro (LLNL) at (415) 422–7503 of their intention to attend at least a

week before the meeting dates to facilitate planning for accommodation.

Minutes of the meeting will be prepared and placed in the NRC Public Document Room. At the end of the peer review process, individual members of the committee will submit their comments to the committee chairman. The chairman will prepare his personal comments in the form of a report to the NRC, and will enclose all comments from the individual members. The review will be done in two phases. In the first phase, the review will be limited to the draft NUREG-1150, and will be completed in December 1987. The changes and improvements to the draft NUREG-1150 that are being performed currently will be reviewed in the second phase and will be completed in July, 1988.

Dated at Rockville, Maryland, this 27th day of August, 1987.

For the Nuclear Regulatory Commission.

R. Wayne Houston,

Acting Director, Division of Reactor Accident Analysis, Office of Nuclear Regulatory Research.

[FR Doc. 87-20219 Filed 8-31-87; 8:45 am]

PACIFIC NORTHWEST ELECTRIC POWER AND CONSERVATION PLANNING COUNCIL

Power Plan; Columbia River Fish and Wildlife Program Amendment

AGENCY: Pacific Northwest Electric Power and Conservation Planning Council.

ACTION: Notice of final amendments.

SUMMARY: On November 15, 1982, pursuant to the Pacific Northwest Electric Power Planning and Conservation Act (Northwest Power Act, 16 U.S.C. 839 et seq.), the Pacific Northwest Electric Power and Conservation Planning Council (Council) adopted a Columbia River Basin Fish and Wildlife Program (Program). The Program has been amended on several occasions since then. On May 29, 1987 certain parties submitted applications to the Council asking that the Program be amended to incorporate certain terms of a settlement agreement regarding fish protection and mitigation measures at Rock Island Dam, located in the State of Washington. In a public meeting on June 10, 1987 the Council authorized issuance of a notice of proposed amendments. The notice was duly published in the Federal Register (52 FR 23223, June 18, 1987). Hearings on the proposed amendments were held in Portland, Oregon on June 23, in Boise, Idaho on

June 30, and in Helena, Montana and Seattle, Washington on July 1, 1987. In addition, a consultation with interested parties was held in Wenatchee, Washington on June 17, 1987, and opportunity for oral and written public comment was provided through July 15, 1987. Written comment was received from five (5) entities and individuals.

Final Amendments

The Council has considered fully the issues and public comment in this rulemaking, and has evaluated the proposed alternatives under the standards for Program measures stated in section 4(h) of the Northwest Power Act. The Council hereby amends the Columbia River Basin Fish and Wildlife Program as follows:

1. In section 402, the second paragraph is amended to read as follows:

In 1982, the Council called for development of mechanical bypass systems at five public utility district dams regulated by FERC in the mid-Columbia area. In 1984, operators of four of the five dams agreed to develop bypass systems as part of a settlement with fish and wildlife agencies and tribes, which had petitioned FERC to make bypass a condition of license renewals for the dams. Spill, which is to be used to protect fish until the bypass systems are operating, is to be shaped in coordination with the fish and wildlife agencies and tribes. At Priest Rapids Dam, the Council called for the study of a short-haul fish transportation program, while a prototype bypass system is being tested at the project. In 1987, the Council amended section 404(a) of the program to incorporate provisions of a settlement agreement concerning fish protection measures at Rock Island Dam. The settlement capped several years of litigation over the advisability of mechanical bypass systems for juvenile fish, whether a hatchery would be a reasonable substitute, what level of spill would be appropriate to protect juvenile fish, and other issues. The settlement agreement calls for the development of juvenile bypass systems, and installation of the systems if certain critieria are satisfied. The agreement also provides for the creation of an innovative "Fisheries Conservation Account," which the joint fishery parties who have signed the agreement may use for bypass studies, bypass development, or to purchase spill. The agreement specifies spill levels and provides for studies of summer spill. A hatchery and satellite facilities will be constructed promptly, and habitat and other studies will be conducted to help determine the proper

use of the fish produced. Changes were

also made in section 604(b)(1) of the program, concerning adult fishway operating criteria and modifications.

2. In section 403, sections 403(a)(2) (A) and (B), and 403(a)(10) are amended to read:

403-Measures

(a) Mid Columbia River Passage

(2) The FERC shall require Chelan County PUD to: (A) Complete testing and evaluation of prototype collection and bypass systems at Rocky Reach and Rock Island dams and report the results of such tests and evaluation to the Council. The evaluation shall compare the effectiveness of the prototype collection and bypass systems with the best available system. If the Council determines that the systems tested at Rocky Reach are not the best available, the FERC shall require the PUD to evaluate alternative collection and bypass systems. Prototype collection and bypass systems for Rock Island dam shall be tested and evaluated in accordance with Section B "Juvenile Fish Bypass Systems" or Section C "Fisheries Conservation Account" of the Settlement dated April 24, 1987, filed in the relicensing proceeding for Project No. 943 and FERC Docket Nos. E-9569,

(B) Complete installation of collection and bypass systems which have been approved by the Council at Rocky Reach Dam. Complete installation of collection and bypass systems at Rock Island Dam in accordance with Section B "Juvenile Fish Bypass Systems" or Section C "Fisheries Conservation Account" of the Settlement Agreement dated April 24, 1987, filed in the relicensing proceeding for Project No. 943 and FERC Docket Nos. E-9569, et al.

(10) The FERC shall require Douglas, Chelan, and Grant County PUDs, in consultation with the fish and wildlife agencies and tribes, to develop plans for spills at their respective projects. These plans shall be developed by March 1 of each year. The FERC shall require the PUDs to use their best efforts to provide spills which will achieve smolt survival comparable to that achievable by the best available collection and bypass systems. In the case of Wells, Rocky Reach, Wanapum and Priest Rapids dams, the FERC shall require the PUDs to provide spills of at least 20 percent of the average daily flow at each project for any 30 out of 60 days when the smolts are present. Such spills may be used during the early nighttime hours for maximum effectiveness and such spill

shall be provided for the period from April 15 through June 15 of each year. During the 30 days when smolts are present, a PUD may be allowed to spill less than 20 percent of the average daily flow only if the PUD can demonstrate to the satisfaction of the Council that at least 90 percent smolt survival at a particular project can be achieved by such reduced spills. In the case of Wells, Rocky Reach, and Wanapum dams, the FERC shall require the operating PUD to implement such plans for spills at each project until a collection and bypass system is in operation. At Priest Rapids Dam, the FERC shall require Grant County PUD to implement such plans until a collection and bypass system is in operation, or until the Council has determined that the short-haul transportation program is likely to be as effective as a collection and bypass system. At Rock Island Dam, FERC shall require spill in accordance with Section C "Fisheries Conservation Account" or Section D "Spill Program" of the Settlement Agreement dated April 24, 1987, filed in the relicensing proceeding for Project No. 943 and Docket Nos. E-9569, et al.

Section 604

- 3. In section 604, section 604(a)(b)(4) is amended to read:
- (b) Operation and Maintenance of Adult Fishways
- (4) At Rock Island Project, FERC shall direct Chelan PUD to implement the operating criteria and adult fishway modifications provided in Section F, "Adult Fish Ladders" of the Settlement Agreement dated April 24, 1987, filed in the relicensing proceeding for Project No. 943 and FERC Docket Nos. E-9569, et al.
- 4. In section 703, section 703(f)(6) is amended to read:

703-Measures

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(f) Construction of Major Hatchery Facilities

(6) FERC shall direct Chelan County PUD to fund design, construction, operation and maintenance of a hatchery program, including satellite facilities, for Rock Island Project in accordance with Section E "Hatchery-Based Compensation" of the Settlement Agreement dated April 24, 1987, filed in the relicensing proceeding for Project No. 943 and Docket Nos. E-9569, et al.

Response to Comments

Summary of Comments

Comments favoring the proposed amendment were received from the Columbia Basin Fish and Wildlife Authority, the Icicle Valley Chapter of Trout Unlimited (Leavenworth, Washington), the Washington Department of Fisheries, and a fisheries biologist of the United States Bureau of Indian Affairs. In addition, at a June 17, 1987 consultation representatives of parties to the settlement agreement expressed support for the proposed amendments. In response to the Council's request, the parties to the settlement agreement indicated that they would cooperate in the Council's system planning process as they design and implement their production facilities. No adverse comments were received.

The Bonneville Power Administration supported the agreement in general terms, and also noted its understanding that the settlement "does not conflict with, and will not abrogate, the mid-Columbia hourly coordination contract, the Pacific Northwest Coordination Agreement, and any other agreements affecting the operation of the Rock Island Project."

Council Response

The Council appreciates these expressions of support and cooperation. The Council agrees with Bonneville's understanding, and does not intend that the Program amendment supplant or modify any existing obligations under other contracts or agreements.

Edward Sheets,

Executive Director,

[FR Doc. 87-20023 Filed 8-31-87; 8:45 am]

SMALL BUSINESS ADMINISTRATION

[Disaster Loan Area #2288]

Declaration of Disaster Loan Area; Illinois

As a result of the President's major disaster declaration on August 21, 1987, I find that the following areas in the State of Illinois constitute a disaster loan area as a result of severe storms and flooding beginning on August 13, 1987: Elk Grove, Hanover, Leyden, Lyons, Maine, Norwood Park, Palatine, Proviso, River Forest, Riverside, Schaumburg and Wheeling Townships in Cook county, and Addison,

Bloomingdale, Downers Grove, Wayne and York Townships in DuPage County.

Eligible persons, firms and organizations may file application for physical damage until the close of business on October 22, 1987, and for economic injury until the close of business on May 23, 1988, at: Disaster Area 2 Office, Small Business Administration, 120 Ralph McGill Blvd., 14th Floor, Atlanta, Georgia 30308, or other locally announced locations.

The interest rates are:

The second second second	(Percent)	
Homeowners with credit avail- able elsewhere	8.000	
Homeowners without credit available elsewhere	4.000	
Businesses with credit avail- able elsewhere	8.000	
Businesses without credit avail- able elsewhere	4.000	
Businesses (EIDL) without credit available elsewhere	4.000	
Other (non-profit organizations including charitable and reli-		
gious organizations)	9.500	

The number assigned to this disaster is 228806 for physical damage and for economic injury the number is 654500.

(Catalog of Federal Domestic Assistance Programs Nos. 59002 and 59008)

Date: August 25, 1987.

Bernard Kulik,

Deputy Associate Administrator for Disaster Assistance.

[FR Doc. 87-19983 Filed 8-31-87; 8:45 am]

[Disaster Loan Area #2286, Amdt. #1]

Declaration of Disaster Loan Area; Minnesota

The above-numbered Declaration (52 FR 30757) is hereby amended in accordance with the Notice of Amendment to the President's declaration, dated August 21, 1987, to include Anoka County in the State of Minnesota because of damage from severe storms, tornadoes and flooding beginning on or about July 20, 1987. All other information remains the same; i.e., the termination date for filing applications for physical damage is the close of business on October 5, 1987, and for economic injury until the close of business on May 6, 1988.

(Catalog of Federal Domestic Assistance Programs Nos. 59002 and 59008) Date: August 25, 1987.

Bernard Kulik.

Deputy Associate Administrator for Disaster Assistance.

[FR Doc. 87-19984 Filed 8-31-87; 8:45 am]

[License No. 06/06-0182]

Surrender of License; First SBIC of Arkansas, Inc.

Notice is hereby given that First SBIC of Arkansas, 700 Worthen Bank Building, Little Rock, Arkansas 72201 has surrendered its License to operate as a small business investment company under the Small Business Investment Act of 1958, as amended (Act). First SBIC of Arkansas was licensed by the Small Business Administration on September 7, 1976. Under the authority vested by the Act and pursuant to the Regulation promulgated thereunder, the surrender was accepted on August 19, 1987, and accordingly, all rights, privileges, and franchises therefrom have been terminated.

(Catalog of Federal Domestic Assistance Program No. 59.011, Small Business Investment Companies)

Robert G. Lineberry,

Deputy Associate Administrator for Investment.

Dated: August 24, 1987.

[FR Doc 87-19985 Filed 8-31-87; 8:45 am]

[License No. 09/12-0075]

Filing of an Application for Exemption Under the Conflict of Interest Regulation; Ritter Partners

Notice is hereby given that Ritter Partners, 3000 Sand Hill Road, Building 2, Suite 215, Menlo Park, California 94025, a Federal Licensee under the Small Business Investment Act of 1958, as amended (the Act), has filed an application with the Small Business Administration (SBA) pursuant to \$ 107.903(b) of the Regulations governing small business investment companies (13 CFR 107.903(b) (1987)) for an exemption from the provisions of the cited Regulation.

The transaction falls within the purview of the cited regulation because Ritter Partners proposes to add a General Partner, Mr. William C. Edwards, 3000 Sand Hill Road, Bldg. 2, Suite 215, Menlo Park, California to the Board of Directors of a portfolio concern, Natural Language, Inc., 1786 Fifth Street, Berkeley, California 94710.

Notice is hereby given that any interested person may, not later than fifteen (15) days from the date of publication of this Notice, submit written comments on the proposed action to the Deputy Associate Administrator for Investment, Small Business Administration, 1441 L Street NW., Washington, DC 20416.

A copy of this Notice will be published in a newspaper of general circulation in the Berkeley California

(Catalog of Federal Domestic Assistance Program No. 59.011, Small Business Investment Companies)

Robert G. Lineberry,

Deputy Associate Administrator for Investment

Dated: August 24, 1987.

[FR Doc. 87–19986 Filed 8–31–87; 8:45 am]

BILLING CODE 8025-01-M

DEPARTMENT OF STATE

[Public Notice CM-8/1105]

Joint Working Party of The U.S. Organization For The International Telegraph and Telephone Consultative Committee (CCITT); Notice of Meeting

The Department of State announces that Study Group JWP of the U.S. Organization for the International Telegraph and Telephone Consultative Committee (CCITT) will meet on September 25, 1987 at the Claremont Hotel, Oakland, California at 8:30 a.m.

The purpose of the meeting will be to review results of the meeting of CCITT Study Group XI and prepare and approve ISDN related U.S. contributions to the upcoming meeting of CCITT Study Group XVII, in Geneva.

Member of the general public may attend the meeting and join in the discussion, subject to the instructions of the Chairman. Admittance of public members will be limited to the seating available. Prior to the meeting, persons who plan to attend should so advise the office of Mr. W.F. Utlaut, NTIA/ITS.D, 325 Broadway, Boulder, CO 80303, telephone (303) 497–3500.

Date: August 20, 1987.

Earl S. Barbely,

Director, Office of Technical Standards and Development; Chairman, U.S. CCITT National Committee.

[FR Doc. 87-20006 Filed 8-31-87; 8:45 am] BILLING CODE 4710-07-M

[Public Notice CM-8/1107]

Closed Meeting; Overseas Security Advisory Council

The Department of State announces a meeting of the U.S. State Department-Overseas Security Advisory Council on Friday, October 2, 1987 at 08:30 a.m. in the Westin Hotel, O'Hare, 6100 River Road, Rosemont, Illinois. Pursuant to section 10 (d) of the Federal Advisory Committee Act and 5 U.S.C. 552b(c)(4), it has been determined the meeting will be closed to the public. Matters relative to privileged commercial information will be discussed. The agenda calls for the discussion of private sector physical security policies, bomb threat statistics, and security programs at sensitive U.S. Government and private sector locations overseas.

Date: August 24, 1987.

Louis Schwartz, Jr.,

Director of the Diplomatic Security Service, and Chairman of the Overseas Security Advisory Council.

[FR Doc. 87-20007 Filed 8-31-87; 8:45 am]

BILLING CODE 4710-24-M

Meeting; Fine Arts Committee

The Fine Arts Committee of the Department of State will meet on Friday, September 25, 1987 at 2:30 p.m. in the John Quincy Adams State Drawing Room. The meeting will last approximately until 4:00 p.m. and is open to the public.

The agenda for the committee meeting will include a summary of the work of the Fine Arts Office since its last meeting in March 1987, the announcement of gifts, loans and financial contributions since January 1, 1987, and a report on the architectural work to be done in the office of the Deputy Secretary of State.

Public access to the Department of State is controlled. Members of the public wishing to take part in the meeting should telephone the Fine Arts Office by Monday, September 21, 1987, telephone (202) 647–1990 to make arrangements to enter the building. The public may take part in the discussion as long as time permits and at the discretion of the chairman.

Date: August 19, 1987.

Clement E. Conger,

Chairman, Fine Arts Committee.

[FR Doc. 87-20008 Filed 8-31-87; 8:45 am]

BILLING CODE 4710-38

DEPARTMENT OF TRANSPORTATION

Office of the Secretary

Reports, Forms, and Recordkeeping Requirements; Submittals to OMB on August 26, 1987

AGENCY: Office of the Secretary, DOT.
ACTION: Notice.

SUMMARY: This notice lists those forms, reports, and recordkeeping requirements imposed upon the public which were transmitted by the Department of Transportation on August 26, 1987, to the Office of Management and Budget (OMB) for its approval in accordance with the requirements of the Paperwork Reduction Act of 1980 (44 U.S.C. Chapter 35).

FOR FURTHER INFORMATION CONTACT:
John Chandler, Annette Wilson, or
Cordelia Shepherd, Information
Requirements Division, M-34, Office of
the Secretary of Transportation, 400
Seventh Street, SW., Washington, DC
20590, telephone (202) 366-4735, or Gary
Waxman or Sam Fairchild, Office of
Management and Budget, New
Executive Office Building, Room 3228,
Washington, DC 20503, (202) 395-7340.

SUPPLEMENTARY INFORMATION:

Background

Section 3507 of Title 44 of the United States Code, as adopted by the Paperwork Reduction Act of 1980, requires that agencies prepare a notice for publication in the Federal Register. listing those information collection requests submitted to the Office of Management and Budget (OMB) for initial, approval, or for renewal under that Act. OMB reviews and approves agency submittals in accordance with criteria set forth in that ACT. In carrying out its responsibilities, OMB also considers public comments on the proposed forms, reporting and recordkeeping requirements. OMB approval of an information collection requirement must be renewed at least once every three years.

Information Availability and Comments

Copies of the DOT information collection requests submitted to OMB may be obtained from the DOT officials listed in the "For Furthr Information Contact" paragraph set forth above. Comments on the requests should be forwarded, as quickly as possible, directly to the OMB officials listed in the "For Further Information Contact" paragraph set forth above. If you anticipate submitting substantive comments, but find that more than 10

days from the date of publication are needed to prepare them, please notify the OMB officials of your intent immediately.

Items Submitted for Review by OMB

The following information collection requests were submitted to OMB on August 26, 1987.

DOT No.: 2967 OMB No.: 2115-0139

Administration: U.S. Coast Guard Title: Hazardous Materials Used as

Ships' Stores Aboard Ships
Need for Information: This information
collection requirement is needed to
regulate the transportation, stowage
and use of ships' stores and dangerous

supplies.

Proposed Use of Information: Coast
Guard uses the information to: (1)
Determine whether a product meets
definitions of hazardous materials and
to properly classify it: (2) make certain
that the instructions on the label are
adequate to protect users on the
vessel from bodily harm; and, (3)
provide proper safeguards in case of
excessive exposure or accident.

Frequency: On occasion
Burden Estimate: 569 hours
Respondents: Manufacturers of
dangerous products used on ships
Form(s): None.

DOT No.: 2968 OMB No.: 2125–0028 Administration: Federal Highway Administration

Title: Highway Performance Monitoring System (HPMS)

Need for Information: For FHWA to obtain information in evaluating the effectiveness of Federal-aid and highway safety programs. Proposed Use of Information: For the

Proposed Use of Information: For the development and implementation of legislation and in resolution of inquiries, including those from Congress.

Frequency: Annually
Burden Estimate: 82,820 hours
Respondents: State highway agencies
Form(s): None

DOT No.: 2969 OMB No.: 2115–0545 Administration: U.S. Coast Guard Title: Deepwater Port Liability Fund Administration

Need for Information: Coast Guard needs this collection of information to administer and manage the Deepwater Port Liability Fund per 33 USC 1517.

Proposed Use of Information: The Coast Guard uses this information to ensure that the owner, operator or guarantor of vessels using deepwater ports have financial responsibility coverage sufficient to meet their pollution liability. It is further used to process and settle any claim made against the "Fund" for cleanup costs or damages resulting from oil pollution.

Frequency: On occasion
Burden Estimate: 143 Hours
Respondents: Owners and operators of
vessels using deepwater ports
Form(s): None

DOT No.: 2970 OMB No.: New Administration: Maritime Administration

Title: Tug Operators Report
Need for Information: Defense/
Emergency Mobilization Planning
Proposed Use of Information: To
determine the characteristics and
availability of tug boats to move
Ready Reserve Force (RRF) cargo
vessels.

Frequency: On occasion
Burden Estimate: 120 hours
Respondents: Tug owners/operators
Form(s): None

DOT No.: 2971
OMB No.: 2120-0001
Administration: Federal Aviation
Administration

Administration

Title: Notice of Proposed (Actual)

Construction or Alteration

Need for Information: This information collection requirement is needed to ensure that any construction or alteration of a structure does not interfere with the safe and efficient use of navigable airspace by aircraft.

Proposed Use of Information: The information is used to establish minimum flight altitudes and procedures, to protect electronic air navigational aids from electromagnetic interference, and provide accurate charting and other notification to airmen.

Frequency: On occasion
Burden Estimate: 15,110 hours
Respondents: All (Anyone proposing to
construct or alter a structure)
Form(s): FAA Form 7460-1, 7460-2,
7460-11

DOT No.: 2972
OMB No.: New
Administration: Office of the Secretary
Title: Drug Testing Control Form
Need for Information: To provide
information needed to establish and

maintain a chain of custody.

Proposed Use of Information: Control form for chain of custody, with affidavit and other pertinent

information.

Frequency: Once per collection

Burden Estimate: 5,050 hours/year @

30,000 samples/year

Respondents: Individuals undergoing testing, collectors, and lab staff Form(s): None

DOT No.: 2973 OMB No.: 2120-0015 Administration: Federal Aviation Administration

Title: FAA Airport Master Record
Need for Information: The information is
needed to keep the Airport Master
Record up to date on the name,
address, phone number and physical
layout of existing airports.

Proposed Use of Information: The data is the basic source of information for private, state and Federal aeronautical charts and publications. Frequency: On occasion/Annually Burden Estimate: 4,000 hours Respondents: State and local governments, businesses (anyone who

has a landing area)
Form(s): FAA Forms 5010-1, -2, -3, -5

Issued in Washington, DC, on August 26, 1987.

Richard B. Chapman,

Acting Director of Information Resource Management.

[FR Doc. 87-20048 Filed 8-31-87; 8:45 am]

Federal Aviation Administration

FAA Approval of Noise Compatibility Program; Tampa International Airport, Tampa, FL

AGENCY: Federal Aviation Administration (FAA), DOT. ACTION: Notice.

SUMMARY: The FAA announces its findings on the noise compatibility program submitted by the Hillsborough County Aviation Authority under the provisions of Title I of the Aviation Safety and Noise Abatement Act (ASNA) of 1979 (Pub. L. 96-193) and 14 CFR Part 150. These findings are made in recognition of the description of Federal and non-Federal responsibilities in Senate Report No. 96-52 (1980). On January 21, 1987, the FAA determined that the noise exposure maps submitted by the Hillsborough County Aviation Authority under Part 150 were in compliance with applicable requirements. On July 15, 1987, the Administrator approved the Tampa International Airport noise compatibility

EFFECTIVE DATE: The effective date of the FAA's approval of the Tampa International Airport noise compatibility program is July 15, 1987.

FOR FURTHER INFORMATION CONTACT: Mr. Tommy J. Pickering, Plans and Programs Metro Area Manager, Federal Aviation Administration, Orlando Airports District Office, 4100 Tradecenter Street, Orlando, Florida 32827, (305) 648–6583. Documents reflecting this FAA action may be obtained from the same individual.

SUPPLEMENTARY INFORMATION: This notice announces that the FAA has given its overall approval to the noise compatibility program for Tampa International Airport, effective July 15, 1987.

Under section 104(a) of the Aviation Safety and Noise Abatement Act (ASNA) of 1979, an airport operator who has previously submitted a noise exposure map may submit to the FAA a noise compatibility program which sets forth the measures taken or proposed by the airport operator for the reduction of existing noncompatible land uses and prevention of additional noncompatible land uses within the area covered by the noise exposure maps. The Act requires such programs to be developed in consultation with interested and affected parties including local communities, government agenices, airport users, and FAA personnel.

Each airport noise compatibility program developed in accordance with the Federal Aviation Regulations (FAR) Part 150 is a local program, not a Federal program. The FAA does not substitute its judgment for that of the airport proprietor with respect to which measures would be recommended for action. The FAA's approval or disapproval of FAR Part 150 program recommendations is measured according to the standards expressed in Part 150 and the Aviation Safety and Noise Abatement Act of 1979, and is limited to the following determinations:

The noise compatibility program was developed in accordance with the provisions and procedures of FAR Part 150:

Program measures are reasonably consistent with achieving the goals of reducing existing noncompatible land uses around the airport and preventing the introduction of additional noncompatible land uses;

Program measures would not create an undue burden on interstate or foreign commerce, unjustly discriminate against types or classes of aeronautical uses, violate the terms of airport grant agreements, or intrude into areas preempted by the Federal Government.

Program measures relating to the use of flight procedures can be implemented within the period covered by the program without derogating safety, adversely affecting the efficient use and management of the Navigable Airspace

and Air Traffic Control Systems, or adversely affecting other powers and responsibilities of the Administrator prescribed by law.

Specific limitations with respect to FAA's approval of the airport noise compatibility program are delineated in FAR Part 150, § 150.5. Approval is not a determination concerning the acceptability of land uses under Federal. State, or local law. Approval does not by itself constitute an FAA implementing action. A request for Federal action or approval to implement specific noise compatibility measures may be required, and an FAA decision on the request may require an environmental assessment of the proposed action. Approval does not constitute a commitment by the FAA to financially assist in the implementation of the program nor a determination that all measures covered by the program are eligible for grant-in-aid funding from the FAA under the Airport and Airway Improvement Act of 1982. Where Federal funding is sought, requests for project grants must be submitted to the FAA Airports District Office in Orlando, Florida.

The Hillsborough County Aviation Authority submitted to the FAA on November 5, 1986, the noise exposure maps, descriptions, and other documentation produced during the noise compatibility planning study conducted from February 8, 1985, through November 4, 1985. The Tampa International Airport noise exposure maps were determined by FAA to be in compliance with applicable requirements on January 21, 1987. Notice of this determination was published in the Federal Register on February 10, 1987.

The Tampa International Airport study contains a proposed noise compatibility program comprised of actions designed for phased implementation by airport management and adjacent jurisdictions from the date of study completion to the year 1992. It was requested that the FAA evaluate and approve this material as a noise compatibility program as described in section 104(b) of the Act. The FAA began its review of the program on January 21, 1987, and was required by a provision of the Act to approve or disapprove the program within 180 days other than the use of new flight procedures for noise control. Failure to approve or disapprove such program within the 180-day period shall be deemed to be an approval of such

The submitted program contained twelve proposed actions for noise

mitigation, on and off the airport. The FAA completed its review and determined that the procedural and substantive requirements of the Act and FAR Part 150 have been satisfied. The overall program, therefore, was approved by the Administrator effective July 15, 1987.

Outright approval was granted for all of the specific program elements. The approval action was for the following

program elements:

A. Measure No. 2/3—Use southerly traffic flows whenever possible

B. Measure No. 13—Encourage operations of turbojet aircraft to use ATA-recommended noise abatement arrival procedures

C. Measure No. 14—Designate engine run-up areas

D. Measure No. 15—Augment vegetation noise barrier along the western perimeter of the airport

E. Measure No. 26/27—Establish a helipad on the east side of the airport

- F. Measure No. 33—Zoning for compatible use
- G. Measure No. 34—Overlay zoning

H. Measure No. 38—Purchase of undeveloped land

- Measure No. 40—Soundproofing new construction (building code amendment)
- J. Measure No. 42—Public information program

K. Measure No. 46—Acquisition of developed property, and

L. Measure No. 47—Purchase of navigation easements.

These determinations are set forth in detail in a Record of Approval endorsed by the Administrator on July 15, 1987. The Record of Approval, as well as other evaluation materials and the documents comprising the submittal, are available for review at the FAA office listed above and at the administrative offices of the Hillsborough County Aviation Authority.

Issued in Orlando, Florida, on August 17, 1987.

James E. Sheppard,

Manager, Orlando Airports District Office. [FR Doc. 87–19994 Filed 8–31–87; 8:45 am] BILLING CODE 4910-13-M

Maritime Administration

Approval of Applicant as Trustee; Bank of Delaware

Notice is hereby given that the Bank of Delaware, Wilmington, Delaware, with offices at 300 Delaware Avenue, Wilmington, Delaware, has been approved as Trustee pursuant to Pub. L. 89–346 and 46 CFR 221.21 through 221.30.

Dated: August 25, 1987.

By Order of the Maritime Administrator. James E. Saari,

Secretary.

[FR Doc. 87-20036 Filed 8-31-87; 8:45 am] BILLING CODE 4901-81-M

Research and Special Programs Administration

[Docket No. 87-3W; Notice 1]

Transportation of Natural and Other Gas by Pipeline; Petition for Waiver; Southern Natural Gas Co.

The Southern Natural Gas Company (Southern) has petitioned the Office of Pipeline Safety for a waiver from compliance with 49 CFR 192.553(d). which prohibits, when uprating a pipeline, the establishment of a maximum allowable operating pressure (MAOP) greater than would be permitted for a new pipeline segment constructed of the same materials in the same location. Anticipating increased demand for gas, Southern has requested a waiver to permit the MAOP of 7 pipeline segments to be uprated from 500 to 650 psig. The alternative is replacement of the segments at a cost of \$1.6 million. The seven segments are all in Class 3 locations, which are generally characterized as areas with 46 or more occupied buildings per mile of pipeline (see § 192.5). The segments are located in St. Clair and Talladega Counties of Alabama, and they are shown on vicinity plat drawing MD-472-1, site plat drawing MD-472, and class location surveys O-NSE-34, 35, and 36, which are available in the docket.

The segments are part of Southern's 24-inch Second North Main Line. Five of the segments were constructed in 1953; the other two were constructed in 1963. The pipe in all segments is 24 inches in diameter, with 0.250-inch wall thickness. It was manufactured to API Standard 5LX, grade 52, and the pipeline was constructed in accordance with the "USAS B31.8 Standard Code for Pressure Piping. Gas Transmission and Distribution systems." The 1953 segments were pressure tested in place with gas to 750 psig, and the 1963 segments were tested with gas to 585

psig.

The MAOP's on the Second North Main Line are 750 and 650 psig, moving west to east (MP 0.0 to MP 231.0) and 500 psig in the remaining 6 miles. They were established in 1970 under § 192.619(a)(3) as the highest actual operating pressure in the preceding five years. The segments in question are located in the last six miles (MP 231.795 to MP 236.646) and comprise a total of 3.4 miles. The construction of these

segments is substantially the same as the rest of the Second North Main Line, which operates at a higher MAOP.

Southern states that the Second North Main Line is coated, and the coating is in good condition. They also state that there have been no leaks or failures, no operating or maintenance problems, and no shorted casings. The line has minimum cover of 36 inches.

The sections of this pipeline that were operated at 650 psig in the five years prior to November 1970 have been able to continue to operate at that pressure in Class 3 locations, after suitable hydrostatic testing under § 192.611(c)(2). Because construction of the entire pipeline is substantially the same, had the 7 segments in question been operated at 650 psig during the 1965—1970 period, they too could be qualified for continued operation at 650 psig after hydrostatic testing.

In summary, the Second North Main Line is 237 miles long. The MAOP is at least 650 psig for 231 miles, including segments in Class 3. Southern plans to uprate the remaining 6 miles to a 650-psig MAOP and has requested that § 192.553(d) be waived to permit them to uprate 3.4 miles of pipeline currently limited by that section.

Because the length of the pipeline segments for which the waiver has been requested is short and because there are similarly constructed pipeline segments already qualified for 650 psig in Class 3 locations along the Second North Main Line, it seems reasonable to waive § 192.553(d) for these segments. There is no reason to anticipate a lesser level of safe performance for the segments for which the waiver has been requested than has been demonstrated by the other similar Class 3 segments. Thus, granting of this waiver would provide uniform treatment for the Second North Main Line taken as a unit and would not impair the safety of that line. In addition, there do not appear to be any unusual risks associated with the populations in proximity to the segments in question. In view of the foregoing, OPS proposes to grant the requested waiver.

Interested parties are invited to comment on the proposed waiver by submitting in duplicate such data, views, or arguments as they may desire.

Comments should identify the Docket and Notice numbers and be submitted to: Dockets Unit, Office of Hazardous Materials Transportation, Research and Special Programs Administration, U.S. Department of Transportation, 400 Seventh Street, SW., Room 8426, Washington, DC 20590.

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All comments received before October 1, 1987 will be considered before final action is taken. Late filed comments will be considered so far as practicable. All comments will be available for inspection at the Dockets Unit, Room 8426, Research and Special Programs Administration, between the hours of 8:30 a.m. and 5:00 p.m., before and after the closing date. No public hearing is contemplated, but one may be held at a time and place set in a Notice in the Federal Register if requested by an interested person desiring to comment at a public hearing and raising a genuine issue.

Issued in Washington, DC, on August 26, 1987.

Richard L. Beam,

Director, Office of Pipeline Safety Research and Special Programs Administration.

[FR Doc. 87-19990 Filed 8-31-87; 8:45 am]

BILLING CODE 4910-60-M

DEPARTMENT OF THE TREASURY

Public Information Collection Requirements Submitted to OMB for Review

Date: August 27, 1987.

The Department of Treasury has submitted the following public information collection requirement(s) to OMB for review and clearance under the Paperwork Reduction Act of 1980, Pub. L. 96–511. Copies of the submission(s) may be obtained by calling the Treasury Bureau Clearance Officer listed. Comments to the OMB reviewer listed and to the Treasury Department Clearance Officer, Department of the Treasury, Room 2224, 15th and Pennsylvania Avenue, NW., Washington, DC 20220.

Internal Revenue Service

OMB number: 1545-0922
Form number: 8329 and 8330
Type of review: Extension
Title: Lender's Information Return for
Mortgage Credit Certificates (MCCs);
Issuer's Quarterly Information Return
for Mortgage Credit Certificates
(MCCs)

Description: These forms will be used by lending institutions who issued qualified indebtedness amounts based on mortgage credit certificates. The information on this form will be matched with the information supplied by the issuers of MCCs.

Respondents: State or local governments, Businesses or other for-

Estimated burden: 27,500 hours Clearance officer: Garrick Shear (202) 535–4297, Internal Revenue Service, Room 5571, 1111 Constitution Avenue, NW., Washington, DC 20224 OMB reviewer: Milo Sunderhauf (202) 395–6880, Office of Management and Budget, Room 3208 New Executive Office Building, Washington, DC 20503

Dale A. Mergan,

Departmental Reports Management Officer. [FR Doc 87–20054 Filed 8–31–87; 8:45 am] BILLING CODE 4819–25-M

Public Information Collection Requirements Submitted to OMB for Review.

Date: August 27, 1987.

The Department of Treasury has made revisions and resubmitted the following public information collection requirement(s) to OMB for review and clearance under the Paperwork Reduction Act of 1980, Pub. L. 96-511. Copies of the submission(s) may be obtained by calling the Treasury Bureau Clearance Officer listed. Comments regarding these information collections should be addressed to the OMB reviewer listed and to the Treasury Department Clearance Officer, Room 2224, Main Treasury Building, 15th and Pennsylvania Avenue, NW., Washington, DC 20220.

Internal Revenue Service

OMB number: 1545–0085
Form number: 1040A
Type of review: Resubmission
Title: U.S. Individual Income Tax Return
Description: This form is used by
individuals to report their income
subject to income tax and to compute
their correct tax liability. The data is
used to verify that the income
reported on the form is correct. The
data is also used for statistical
purposes.

Respondents: Individuals or households Estimated burden: 21,805,522 hours

OMB number: 1545-0099

Form number: 1065 and Schedules D and K-1

Type of review: Resubmission
Title: U.S. Partnership Return of Income,
Capital Gains and Losses, Partner's
Share of Income, Credits, Deductions,
etc.

Description: Internal Revenue Code section 6031 requires partnerships to file returns that show gross income items allowable deductions, partners' names, addresses, and distribution shares, and other information. This information is used to verify correct reporting of partnership items and for general statistics.

Respondents: Individuals or households, Farms, Businesses or other for-profit Estimated burden: 17,930,209 hours OMB number: 1545–0675 Form number: 1040EZ

Type of review: Resubmission
Title: Income Tax Return for Single
Filers with No Dependents

Description: This form is used by certain single individuals to report their income subject to income tax and to compute their correct tax liability. The data is also used to verify that the items reported on the form are correct and are also for general statistics use. Respondents: Individuals or households

Estimated burden: 9,152,703 hours
Clearance Officer: Garrick Shear (202)
535–4297, Room 5571, 1111
Constitution Avenue, NW.,
Washington, DC 20224

OMB Reviewer: Milo Sunderhauf (202) 395–6880, Office of Management and Budget, Room 3208, New Executive Office Building, Washington, DC 20503

Dale A. Morgan,

Department Reports Management Officer. [FR Doc. 87-20055 Filed 8-31-87; 8:45 am] BILLING CODE 4810-25-M

VETERANS ADMINISTRATION

Agency Form Under OMB Review

AGENCY: Veterans Administration. ACTION: Notice.

The Veterans Administration has submitted to OMB for review the following proposal for the collection of information under the provisions of the Paperwork Reduction Act (44 U.S.C. Chapter 35). This document contains a reinstatement and lists the following information: (1) The department or staff office issuing the form, (2) the title of the form, (3) the agency form number, if applicable, (4) a description of the need and its use, (5) how often the form must be filled out, (6) who will be required or asked to report, (7) an estimate of the number of responses, (8) an estimate of the total number of hours needed to fill out the form, and (9) an indication of whether section 3504(h) of Pub. L. 96-511 applies.

ADDRESSES: Copies of the forms and supporting documents may be obtained from Patti Viers, Agency Clearance Officer (732), Veterans Administration, 810 Vermont Avenue, NW., Washington, DC 20420, (202) 233–2146. Comments and questions about the items on the list should be directed to the VA's OMB Desk Officer, Joseph Lackey, Office of Management and Budget, 726 Jackson Place, NW., Washington, DC 20503, (202) 395–7316.

DATES: Comments on the information collection should be directed to the

OMB Desk Officer on or before November 2, 1987.

Dated: August 26, 1987.

By direction of the Administrator.

David A. Cox, Associate Deputy Administrator for

Management. Reinstatement

- 1. Board of Veterans Appeals
- 2. Statement of Accredited
 Representative in Appealed Cases
- 3. VA Form 1-646
- 4. This form is used to ensure that the appellant's rights are protected. This information is used in the final stages of the claims process or appellate review.
- 5. On occasion
- 6. Individuals or households
- 7. 36,975 responses
- 8. 36,975 hours
- 9. Not applicable

[FR Doc. 87-20039 Filed 8-31-87; 8:45 am]
BILLING CODE 8320-01-M

Meeting; Career Development Committee

The Veterans Administration gives notice under Pub. L. 92-463 that a meeting of the Career Development Committee, authorized by 38 U.S.C. 4101, will be held in the State Room of the Governor's House, Rhode Island Avenue at 17th Street, NW., Washington, DC, October 4 through 6, 1987, starting at 7 p.m., October 4. The meeting will be for the purpose of scientific review of applications for appointment to the Career Development Program in the Veterans Administration. The committee advises the Director. Medical Research Service on selection and appointment of Associate Investigators, Research Associates, Clinical Investigators, Medical Investigators, and Senior Medical Investigators.

The meeting will be open to the public up to the seating capacity of the room

from 7 p.m. to 7:30 p.m. on October 4, 1987, to discuss the general status of the program. Because of the limited seating capacity of the room, those who plan to attend should contact Mr. David D. Thomas, Executive Secretary of the Career Development Committee (151]), Veterans Administration Central Office, Washington, DC 20420 (Phone 202–233–2317) prior to September 30, 1987.

The meeting will be closed from 7:30 p.m. to 10 p.m., October 4 and from 8 a.m. to 5 p.m. on October 5 and October 6, for consideration of individual applications for positions in the Career Development Program. This necessarily requires examination of personnel files and discussion and evaluation of the qualification, competence, and potential of the several candidates, disclosure of which would constitute a clearly unwarranted invasion of personal privacy. Accordingly, closure of this portion of the meeting is permitted by section 10(d) of Pub. L. 92-463 as amended, in accordance with subsection (c)(6), 5 U.S.C. 552b.

Minutes of the meeting and rosters of the committee members may be obtained from Mr. David D. Thomas, Chief, Career Development Program, Medical Research Service (151J), Veterans Administration, Washington, DC 20420 (Phone 202–233–2317).

Dated: August 25, 1987.

By direction of the Administrator.

Rosa Marie Fontanez,

Committee Management Officer.
[FR Doc. 87–20040 Filed 8–31–87; 8:45 am]
BILLING CODE 8320-01-M

Meeting; Cooperative Studies Evaluation Committee

The Veterans Administration gives notice under Pub. L. 92–463 that a meeting of the Cooperative Studies Evaluation Committee, authorized by 38 U.S.C. 4101, will be held at the Vista International Hotel, 1400 M Street, NW., Washington, DC, on October 22, 1987.

The meeting will be for the purpose of reviewing proposed cooperative studies and advising the Veterans
Administration on the relevance and feasibility of the studies, the adequacy of the protocols, and the scientific validity and propriety of technical details, including protection of human subjects. The Committee advises the Director, Medical Research Service, through the Chief of the Cooperative Studies Program on its findings.

The meeting will be open to the public up to the seating capacity of the room from 7:30 a.m. to 8 a.m. on October 22, 1987, to discuss the general status of the program. To assure adequate accommodations, those who plan to attend should contact Dr. Ping Huang, Coordinator, Cooperative Studies Evaluation Committee, Veterans Administration Central Office, Washington, DC (202–233–2861), prior to October 8, 1987.

The meeting will be closed from 8 a.m. to 3:30 p.m. on October 22, for consideration of specific proposals in accordance with provisions set forth in subsection 10(d) of Pub. L. 92-463, as amended by Pub. L. 94-409, and as cited in 5 U.S.C. 552b(c)(6) and (9)(B). During this portion of the meeting discussions and decisions will deal with qualifications of personnel conducting the studies and the medical records of patients who are study subjects, the disclosure of which would constitute clearly unwarranted invasion of personal privacy. Additionally, premature disclosure of the Committee's recommendations would likely frustrate implementation of final proposed actions.

Dated: August 25, 1987.

By direction of the Administrator.

Rosa Maria Fontanez,

Committee Management Officer. [FR Doc. 87-20041 Filed 8-31-87; 8:45 am] BILLING CODE 8320-01-M

Sunshine Act Meetings

Federal Register

Vol. 52, No. 169

Tuesday, September 1, 1987

This section of the FEDERAL REGISTER contains notices of meetings published under the "Government in the Sunshine Act" (Pub. L. 94-409) 5 U.S.C. 552b(e)(3).

BOARD OF GOVERNORS OF THE FEDERAL RESERVE SYSTEM

TIME AND DATE: 11:00 a.m., Tuesday, September 8, 1987.

PLACE: Marriner S. Eccles Federal Reserve Board Building, C Street entrance between 20th and 21st Streets, NW., Washington, DC 20551.

STATUS: Closed.

MATTERS TO BE CONSIDERED:

1. Personnel actions (appointments, promotions, assignments, reassignments, and salary actions) involving individual Federal Reserve System employees.

Any items carried forward from a previously announced meeting.

CONTACT PERSON FOR MORE INFORMATION: Mr. Joseph R. Coyne, Assistant to the Board; (202) 452–3204. You may call (202) 452–3207, beginning at approximately 5 p.m. two business days before this meeting, for a recorded announcement of bank and bank holding company applications scheduled for the meeting.

Date: August 28, 1987.

James McAfee,

Associate Secretary of the Board. [FR Doc. 87–20213 Filed 8–28–87; 3:59 pm] BILLING CODE 8210–01-M

FEDERAL MINE SAFETY AND HEALTH REVIEW COMMISSION

August 26, 1987.

TIME AND DATE: 10:00 a.m., Wednesday, September 2, 1987.

PLACE: Room 600, 1730 K Street, NW., Washington, DC.

STATUS: Open.

MATTERS TO BE CONSIDERED: The Commission will consider and act upon the following:

1. Secretary of Labor on behalf of Bushnell v. Cannelton Industries, Inc., Docket No. WEVA 85–273–D. (Issues include whether the judge erred in finding the operator discriminated against the complainant for engaging in rights protected by section 105(c)(1) of the Mine Act, 30 U.S.C. 815(c)(1).)

2. Martha Perando v. Mettiki Coal Corporation, Docket No. YORK 85-12-D. (Issues include whether the judge erred in finding that the operator disciminated against the complainant in violation of section 105(c)(1) of the Mine Act. 30 CFR part 815(c)(1).)

Any person intending to attend this meeting who requires special accessibility features and/or auxiliary aids, such as sign language interpreters, must inform the Commission in advance of those needs. Subject to 20 CFR 2706.150(a)(3) and 2706.160(e).

CONTACT PERSON FOR MORE INFO: Jean Ellen (202) 653–5629.

Jean H. Ellen, Agenda Clerk.

[FR Doc. 87-20083 Filed 8-28-87; 10:34 am] BILLING CODE 6735-01-M

Corrections

Federal Register

Vol. 52, No. 169

Tuesday, September 1, 1987

This section of the FEDERAL REGISTER contains editorial corrections of previously published Presidential, Rule, Proposed Rule, and Notice documents and volumes of the Code of Federal Regulations. These corrections are prepared by the Office of the Federal Register. Agency prepared corrections are issued as signed documents and appear in the appropriate document categories elsewhere in the issue.

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Parts 795 and 799

[OPTS-42076A; FRL-3213-5]

Anthraquinone; Final Reporting and Recordkeeping Requirements and Test Rule

Correction

In rule document 87-12724 beginning on page 21018 in the issue of Thursday, June 4, 1987, make the following corrections:

§ 795.45 [Corrected]

1. On page 21027, in § 795.45(b)(1)(ii), in Table 1, in the first entry of the first column, "14-" should read "4-"

§ 799.500 [Corrected]

2. On page 21030, in the first column, in § 799.500(d)(2)(i), in the 28th line, "(b)" should read "(B)".

BILLING CODE 1505-01-D

ENVIRONMENTAL PROTECTION

40 CFR Part 799

[OPTS-42089; FRL-3221-7]

Testing Consent Order on 3,4-Dichlorobenzotrifluoride and Response to the Interagency Testing Committee

Correction

In rule document 87-14231 beginning on page 23547 in the issue of Tuesday, June 23, 1987, make the following corrections on that page:

 In the second column, under III. Use and Exposure, in the third paragraph, in the fourth line, "Rone-Poulence" should read "Rhone-Poulenc".

2. In the third column, in the first complete paragraph, in the eighth line, "Manufactures" should read "Manufacturers".

BILLING CODE 1505-01-D

DEPARTMENT OF THE INTERIOR

Bureau of Land Management

43 CFR Public Land Order 6653

[NM-940-07-4220-10; NM NM 52805]

Partial Revocation of Public Land Order No. 6525; New Mexico

Correction

In rule document 87-18041 beginning on page 29525 in the issue of Monday, August 10, 1987, make the following correction:

On page 29526, in the first column, in paragraph 3., in the first line, "1981" should read "1987".

BILLING CODE 1505-01-D

DEPARTMENT OF THE INTERIOR

Bureau of Land Management

[NM-940-07-4220-11; NM NM 023643]

Continuation of Withdrawal and Reservation of Lands; New Mexico

Correction

In notice document 87-15026 beginning on page 25087 in the issue of Thursday, July 2, 1987, make the following

On page 25088, in the first column, the fifth line should read "T. 10 S., R. 18W.,".

BILLING CODE 1505-01-D



Tuesday September 1, 1987

Part II

Department of Defense

Office of the Secretary

32 CFR Part 199

Civilian Health and Medical Program of the Uniformed Services (CHAMPUS); Implementation of CHAMPUS DRG-Based Payment System; Final Rule

DEPARTMENT OF DEFENSE

Office of the Secretary

32 CFR Part 199

[DoD Regulation 6010.8-R]

Civilian Health and Medical Program of the Uniformed Services (CHAMPUS); Implementation of a CHAMPUS DRG-Based Payment System

AGENCY: Office of the Secretary, DoD.
ACTION: Final rule.

SUMMARY: In FR Doc. 77–7834, appearing in the Federal Register on April 4, 1977, (42 FR 17972), the Office of the Secretary of Defense published its regulation, DoD 6010.8–R, "Implementation of the Civilian Health and Medical Program of the Uniformed Services (CHAMPUS)," as Part 199 of this title. DoD Regulation 6010.8–R was reissued in the Federal Register on July 1, 1986 (51 FR 24008).

This final rule amends the comprehensive CHAMPUS regulations. DoD 6010.8-R (32 CFR Part 199), pertaining to payment for inpatient hospital services. This final rule implements a DRG-based payment system, which is modeled on the Medicare Prospective Payment System. This final rule also revises the costsharing requirements for beneficiaries other than dependents of active duty members. This cost-sharing change is necessary under a DRG-based payment system to ensure that cost-sharing amounts are equitable. This final rule also establishes an admission and quality review system for CHAMPUS inpatient hospital claims.

EFFECTIVE DATE: October 1, 1987. This final rule is effective for inpatient hospital admissions occurring on or after that date.

FOR FURTHER INFORMATION CONTACT: Stephen E. Isaacson, Policy Branch, OCHAMPUS, telephone (303) 361–4005.

SUPPLEMENTARY INFORMATION: On June 3, 1987, we published a proposed rule to implement a CHAMPUS DRG-based payment system. This rule proposed to change the method of payment for inpatient hospital services under CHAMPUS from a billed charge, retrospective system to a prospective payment system based on diagnosis-related groups (DRGs). We refer the reader to the proposed rule for more detailed explanations of the proposed changes to the reimbursement

procedures and the implementing regulations in 32 CFR Part 199.

We provided a 30-day comment period of the proposed rule. This final rule announces our decisions on the issues raised by commenters in response to our proposed rule.

To assist the reader in reviewing this document, we are providing the Table of Contents below.

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II. Background

A. CHAMPUS Reimbursement—Current Procedures

B. Summary of Legislation

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I. Synopsis

A. Background

Paying on the basis of a fixed, prospective rate, appropriate to the particular diagnosis involved, has been shown to be an equitable, effective method of paying for hospital care. Instead of paying on the basis of billed charges, CHAMPUS is now implementing a Diagnosis Related Groups (DRG) based payment system which will assure fair payments to hospitals, reduce cost sharing requirements for beneficiaries and the government and provide for new procedures to monitor the quality of care.

B. Improvements Upon the Proposed Rule

Based on public comments, several changes are being made to improve upon the proposed rule, issued in June. Among these is an exemption for children's hospitals until a method can be incorporated that will better reflect the unique kind of care these specialty hospitals provide. Another change is authority to exempt states, such as Maryland, that are exempted by Medicare and have equally effective payment methods. Further, to assure reasonable payment amounts to hospitals in areas of generally higher costs, a wage adjustment and an urban/ rural differentiation, as is used under Medicare, will be made. Also, the beneficiary cost-sharing provisions have been revised to assure that no beneficiary will pay more under the new system that under the old method of paying billed charges, and most beneficiaries will pay much less.

C. CHAMPUS DRG System Modeled After Medicare's

Established System Consistent with the Congressional intent, the proposed CHAMPUS system is modeled closely on the Medicare system. The similarities are exemplified by the fact that even though the CHAMPUS population is younger the average CHAMPUS payment amounts are roughly equal to those under Medicare. The costs of capital and indirect and direct medical education will receive the same special

treatment under CHAMPUS that they receive under Medicare. Like Medicare, long-stay or unusually costly cases will receive additional outlier payments. To assure fair payments, amounts have been calculated on the basis of actual CHAMPUS hospital claims during a recent 12-month period.

D. Fiscal Year 1988 Implementation on Schedule

Four years after Medicare's similar system was adopted, CHAMPUS is now ready to proceed with its DRG-based payment system. This interim period has given hospitals time to adjust to the DRG-based system under the government program that is some 40 times larger than CHAMPUS. During this time, CHAMPUS tested a DRG system in South Carolina and learned valuable lessons. CHAMPUS also had the opportunity to compile and analyze extensive data relating exclusively to charges for care under CHAMPUS. Also, the CHAMPUS fiscal intermediaries have had time to put in place the systems to assure smooth administration under the new payment method. In view of all of these activities, there is no need for CHAMPUS to slow down implementation. Nor is there a need to phase in national rates as Medicare did. Phasing is unnecessary because hospital operations have adjusted to the DRG payment method, now fully implemented under Medicare. and CHAMPUS, unlike Medicare, is typically a very small portion of the hospital's income.

E. Reduced Cost Shares for Beneficiaries

From the beneficiary's standpoint, the new CHAMPUS payment system will have a very positive impact. By reducing the payment amount for hospitals, the 25% cost share retired members and their dependents must pay will now be applied to a much lower amount. As a result, the average cost share per hospital stay will be reduced from about \$1135 to about \$760. To assure that cost sharing fairly reflected the value of hospital services provided to the beneficiary, the proposed rule included a new method of calculating cost sharing based on a per deim amount. The final rule continues this more equitable method, and makes a further adjustment: beneficiary cost shares will be the lesser of the per diem calculation method (the new system) or 25% of billed charges (the old system). This assures that while most beneficiaries will pay less under the new system than the old, additionally, no beneficiaries will pay more.

F. Assuring the Quality of Care

To assist in assuring the quality, reasonableness, and appropriateness of care provided CHAMPUS beneficiaries under the DRG based payment system, a new quality review requirement is being established. CHAMPUS is pursuing appropriate arrangements with the Department of Health and Human Services to undertake this important activity in conjunction with current Peer Review Organization program under Medicare. This will enable CHAMPUS to join HCFA In guarding against premature discharges and effectively monitoring the quality of care.

G. Conclusion

In accord with Congressional intent, spiraling costs will be curtailed by a CHAMPUS DRG-based payment system modeled very closely on the Medicare system. With improvements upon the proposed rule, the final rule provides for fair payments to hospitals and reduced outlays and new quality monitoring for beneficiaries. This payment system, carefully developed, will be fully implemented October 1, 1987.

II. Background

A. CHAMPUS Reimbursement—Current Procedures

Paragraph 199.6(e) of DoD 6010.8-R provides for reimbursement of hospitals and skilled nursing facilities on the basis of billed charges/set rates, costrelated reimbursement similar to that used under Title XVIII of the Social Security Act (Medicare), or prospective reimbursement. CHAMPUS has traditionally reimbursed these providers of care based on the providers' billed charges. Largely because of these procedures, CHAMPUS has been subject to rapidly increasing costs, far in excess of the general rate of inflation. This resulted not only from increases in hospitals' charges, but also from the shifting of costs as other third-party payers implemented cost-controlling reimbursement procedures.

B. Summary of Legislation

1. Department of Defense Authorization Act, 1984

The Department of Defense
Authorization Act, 1984, Pub. L. 98–94
amended Title 10, section 1079(j)(2)(A)
of the United States Code and provided
CHAMPUS with the statutory authority
to reimburse institutional providers
based on diagnosis-related groups
(DRGs). Specifically, it provides that
payments "shall be determined to the
extent practicable in accordance with
the same reimbursement rules as apply

to payments to providers of services of the same type under title XVIII of the Social Security Act."

Consolidated Omnibus Budget Reconciliation Act, 1986

On April 7, 1986, the President signed the Consolidated Omnibus Budget Reconciliation Act, Pub. L. 99-272, which contained a provision requiring hospitals which participate in Medicare also to participate in CHAMPUS for inpatient services (see Section 1866(a)(1)(J) of the Social Security Act, 42 U.S.C. 1395cc(a)(1)(J)). Because of questions regarding the effect of this provision, the effective date in section 9122(b) of Pub. L. 99-272, which enacted section 1866(a)(1)(J), was amended by Pub. L. 99-514, section 1895(b)(6), which was signed by the President on October 22, 1986. Section 1866(a)(1)(J) requires all providers participating in Medicare also to participate in CHAMPUS for inpatient hospital services provided pursuant to admissions to hospitals occurring on or after January 1, 1987.

C. Summary of Proposed Amendment to Rule

In the proposed amendment to rule, we set forth new regulations for the CHAMPUS DRG-based payment system intended to apply to inpatient hospital admissions occurring on or after October 1, 1987. We described how the CHAMPUS system is modeled after the Medicare Prospective Payment System: the applicability of the system, both in terms of services and of hospitals affected; how the DRG weighting factors are calculated; how the adjusted standardized amount is calculated; how adjustments for capital and medical education costs are to be made; and how unusual cases (outliers) are to be handled. We also described changes to the cost-sharing requirements for beneficiaries other than dependents of active duty members. Additionally, we set forth procedures for an admission and quality review system for CHAMPUS inpatient hospital claims.

D. Number and Types of Public Comment

We received a total of 34 individual comments which raised a number of issues. The types and volume of commenters were as follows:

- -Hospital Associations-11
- -Hospitals-12
- -Medical Associations-5
- -Medical Review Organizations-3
- -Third-party payers-1
- —State Governments and Organizations—2

We received a number of general comments which do not relate to any particular provision of the CHAMPUS DRG-based payment system, but relate to the system as a whole and our implementation plans. We will address those comments here before we respond to the comments regarding specific sections of our proposed rule.

A number of commenters suggested we delay implementation of the DRG system until at least October 1988 in order to allow hospitals time to prepare for the changes and to ensure that OCHAMPUS has adequate time to develop a comprehensive database and to consider the public comments.

We think delay in the implementation of the DRG system is unnecessary for two main reasons: first, hospitals have already adjusted to prospective payment under Medicare, and second, OCHAMPUS has developed a comprehensive database with which to develop final weights and rates.

We think that hospitals have adjusted to prospective payment under the Medicare system. As a result, the inclusion of prospective payment for CHAMPUS patients should have a minimal effect on hospital operations. The CHAMPUS prospective payment system merely will result in a marginal increase in workload for existing prospective payment-related activities in hospitals such as medical records, billing, and utilization review activities already conducted for Medicare patients. For many hospitals, the net effect of CHAMPUS DRGs on hospital operations will be positive. Instead of managing two radically different payment systems for government beneficiaries (i.e., Medicare and CHAMPUS), in many ways hospitals will have an opportunity to consolidate patient management, billing, and medical records activities to include both CHAMPUS and Medicare patients.

We disagree with the assertion that CHAMPUS has not had adequate time to develop a comprehensive database. The database used for final weight and rate calculations includes over 300,000 records of all CHAMPUS claims processed from July 1, 1986 to June 30, 1987. This represents a full-year sample of claims from the most recent period. We think that this database will provide the most accurate representation of CHAMPUS utilization. The required calculation methods necessary to derive weights and rates have been developed over the past year and tested on a smaller database. Final calculations merely required the application of these methods to the larger database.

In addition to being unnecessary, delaying implementation of the DRGbased payment system would contravene clear Congressional intent that CHAMPUS adopt a payment method similar to that used by Medicare, an intent repeatedly and consistently expressed by Congress since Medicare's system was adopted four years ago.

Some commenters suggested that 30 days was not sufficient time to comment on the proposed rule. The comment period conforms to statutory requirements. Because of years of experience with the Medicare system. the matters discussed in the proposed rule are quite familiar to the health care community. Thus, a 30-day comment period was quite adequate to permit interested parties to consider the proposed rule and provide substantive comments. The many detailed and thorough comments we actually received confirm this. Also, we received no indication that additional comments would have been received, or additional issues raised, had the comment period been longer.

Some commenters mistakenly believe that CHAMPUS will not pay the full DRG-based amount in those cases where a hospital's charges are less than the DRG-based payment amount. Since this is a basic tenet of DRG-based reimbursement, we want to be sure that this misunderstanding is clarified. Except for those few cases which are classified as short-stay outliers, a hospital will always receive the full DRG-based payment regardless of its costs or charges.

In the proposed rule we stated that "we anticipate few, if any, changes in hospital operations as a result of our implementation of the CHAMPUS DRGbased payment system." One commenter stated that the DRG-based payment system would, indeed, affect operations, since hospitals would need more extensive internal concurrent review, additional discharge planning, and more emphasis on claims coding. We recognize these activities may increase as a result of our DRG-based payment system, but they are not required by the system. Rather they are a reaction to the system, and, more importantly, will help to ensure hospital efficiency, which is a major goal of the existing Medicare system and the new CHAMPUS system.

Several commenters expressed a concern that there could be a disruption in services to CHAMPUS patients if CHAMPUS is unable to begin the timely processing of claims on October 1. A primary concern of OCHAMPUS is that we want a smooth transition to DRG-based payments with no adverse impact on providers or benefiticiaries.

We think that all CHAMPUS fiscal intermediaries are adequately prepared to process CHAMPUS claims under DRGs. Since CHAMPUS began exploring the potential for a prospective payment system, the CHAMPUS fiscal intermediaries have been consulted periodically regarding the impact of prospective payment on their current claims payment procedures. Moreover, all but one of the CHAMPUS fiscal intermediaries currently process Medicare claims, making the transition to CHAMPUS prospective payment relatively simple. And finally, since the release of the proposed rule, CHAMPUS has submitted preliminary weights, rates, cutoff thresholds, and other necessary payment procedures to the intermediaries so that they could begin revision of their payment systems. In short, we and the CHAMPUS fiscal intermediaries are confident that there will be a smooth transition from payment based on billed charges to

One commenter suggested we allow our fiscal intermediaries to automate remittance schedules on a weekly basis rather than issuing individual checks and explanation of benefits for every claim. Under current procedures, the fiscal intermediary can combine multiple claims from the same hospital into a single remittance. This procedure will not change under the DRG-based payment system and should resolve the concern of the commenter.

Below we briefly summarize each of the major provisions of the proposed amendment of rule on which we received comments and provide an analysis of the comments and our responses. We have also provided a reference to each provision's designation in the proposed rule. Section VI of this preamble summarizes the changes we are making to the regulation as a result of the comments we received.

III. General Description of CHAMPUS DRG-Based Payment System

A. Modeled on Medicare's Prospective Payment System (PPS)

1. DRGs Used (§ 199.14(a)(1)(i)(A))

The CHAMPUS DRG-based payment system will use the same DRGs used in the most recently available grouper for the Medicare PPS.

Comment—The age breakdown in the Medicare DRG system is only shown as 17 and below. For children's hospitals it is necessary to have a more specific breakdown in ages, as the difference in levels of care between a one-year-old and a 12-year-old will vary significantly.

Response—We realize this problem may be acute for children's hospitals

and, as Medicare, we have decided to exempt children's hospitals from the CHAMPUS DRG system at this time. We intend to study the issue further and bring these hospitals into the system at a later date.

2. Assignment of Discharges to DRGs (§ 199.14(a)(1)(i)(B))

CHAMPUS will use the Health Care Financing Administration (HCFA) FY 1987 "Grouper" program to classify specific hospital discharges within DRGs, so that each discharge is assigned to a DRG based on the patient's age, sex, principal diagnosis, secondary diagnoses, procedures performed and discharge status. In addition, when the discharge data submitted by a hospital show a surgical procedure unrelated to a patient's principal diagnosis, the bill will be returned to the hospital for validation and verification.

Comment—The CHAMPUS DRGbased payment system should always use the Grouper program which is currently used under the PPS.

Response—In general, CHAMPUS intends to use the most recently available HCFA Grouper program. Currently, the most recently available program is the FY 1987 Grouper which we are using for FY 1988. For future years, HCFA revisions to its Grouper will be adopted by CHAMPUS as soon as practicable following availability of the Grouper.

Comment—It is unnecessary to verify the diagnosis and procedure codes for claims grouped into DRG 468 (unrelated operating room procedure), especially since these claims are later reviewed by the Peer Review Organization (PRO).

Response—We do not believe this is unnecessary. Since, by definition, there is something unusual about the claim data for a DRG 468, we believe it is prudent to verify the data initially to avoid payment errors. However, we have modified this requirement to make it less of a burden on hospitals. We will require only that the fiscal intermediary review such claims.

B. Beneficiary Eligibility

If a beneficiary is eligible for CHAMPUS coverage during any part of his/her inpatient confinement, the claim shall be processed as if the beneficiary were eligible for the entire stay. The beneficiary's cost-sharing status is to be determined based on his/her sponsor's status at the time of admission.

Comment—The rule should specify whether the fiscal intermediary of the PRO is responsible for determining a beneficiary's status.

Response—This determination is always the responsibility of the FI and must be made prior to payment of the claim.

C. Basis of Payment

Payment on a Per Discharge Basis (§ 199.14(a)(1)(i)(C)(2))

Under the CHAMPUS DRG-based payment system, hospitals are paid a predetermined amount per discharge for inpatient hospital services.

Comment—OCHAMPUS should have a provision which permits interim payments on long lengths-of-stay.

Response—In general, we have modeled the CHAMPUS DRG-based payment system on the Medicare system. Medicare intends to eliminate interim payments on long stay outliers. Therefore, we intend to follow this policy.

2. Discharges and Transfers

a. Discharges

(§ 199.14(a)(1)(i)(C)(6)(i)). In this section we listed those actions which qualify as discharges and are eligible for full DRG-based payment.

Comment—A transfer from the care of a hospital included under the CHAMPUS DRG-based payment system to a hospital or unit that is excluded from the system is classified as a transfer, but it should be a discharge.

Response—We agree. This is a discharge under the Medicare PPS, and this was an error in our proposed rule. We have made the change to classify such actions as discharges.

b. Payment to a hospital transferring an inpatient to another hospital (§ 199.14(a)(1)(i)(C)(6)(iv)). In the case of a transfer, the transferring hospital is to be paid a per diem rate not to exceed the DRG-based payment that would have been paid if the patient had been discharged to another setting.

Comment—The method of reimbursement for transfers has not been addressed in a manner to assure adequate payment in the case of multiple transfers of a patient among various hospitals.

Response—We have revised the language to specifically indicate that transferring hospitals can receive additional payment for cases which meet the criteria for long-stay or cost outliners.

Comment—Transfers to military treatment facilities (MTFs) should be classified as discharges rather than as transfers.

Response—MTFs have the primary responsibility and statutory requirement to provide care to CHAMPUS

beneficiaries when space is available. Transfers to hospitals excluded from the CHAMPUS DRG-based payment system are considered discharges because the transferred patient receives subsequent care in a facility that is organized for treatment of conditions distinctly unlike treatment provided in the non-exempt acute care facility. In the case of a transfer to an MTF, however, the patient will receive continuing care comparable to that received in the non-exempt acute care transferring hospital. Therefore, transfers of this nature will be treated just as if the hospital transferred the patient to another non-exempt acute care facility. That is, the transferring hospital will be paid a DRG-specific per diem amount.

Comment—Currently, emergency room physicians must contact the MTF for permission to admit all patients in distress, but not in a life threatening status. Will there be any consideration given to reducing these emergency room transfers under DRG payment?

Response—All CHAMPUS
beneficiaries who live within catchment
areas of MTFs are required to first seek
inpatient care in the MTF before going
to a civilian hospital. That is the purpose
of the requirement cited in this
comment, and it will not change as a
result of DRG-based reimbursement.
Thus, if the MTF can treat the patient,
the patient will be required to go to the
MTF for inpatient services, and the
civilian hospital will be reimbursed only
for the emergency room services on an
outpatient basis.

3. Applicability of the DRG System

a. Areas affected
(§ 199.14(a)(1)(ii)(A)). The CHAMPUS
DRG-based payment system shall apply
to hospitals' services in the fifty states,
the District of Columbia, and Puerto
Rico. There are no exemptions for
services in states which have
implemented a separate DRG-based
payment system or similar payment
system in order to control costs.

Comment—The proposed rule states that the CHAMPUS DRG-based payment system will be applicable to the six CHAMPUS Reform Initiative (CRI) demonstration states. The CRI request for proposals allows contractors to use their own reimbursement system for CHAMPUS Prime.

Response—DRG-based reimbursement will still be required in the demonstration states for all claims which are not CHAMPUS Prime, as it will apply to all standard CHAMPUS claims nationwide. Within CHAMPUS Prime, contractors can use the CHAMPUS DRG system or any other

reimbursement system, subject to the requirements of the CRI contracts.

Comment—Any state payment system granted a federal waiver for the purposes of Medicare reimbursement should be automatically exempt from the CHAMPUS system.

Response-The intent of not granting waivers to individual states was to ensure uniformity of payments throughout the country and to ensure that payments in all states would be adequately controlled. The comments we received argued persuasively that the cost controls in those states exempt from the Medicare PPS are adequate to ensure savings comparable to the CHAMPUS DRG-based payment system. Moreover, it would be disruptive if a single payer, particularly a major Federal payer, chose to exempt their beneficiaries from the state system. As a result, we have revised our position to allow states to be exempt from the CHAMPUS DRG-based payment system under the following circumstances:

The State must be exempt from the Medicare PPS:

The State must request, in writing, that it be exempt from the CHAMPUS DRG-based payment system; and

3. Payments in the State must continue to be at a level to maintain savings comparable to those which would be achieved under the CHAMPUS DRGbased payment system. This exemption is based on savings achieved under Medicare, but because of the differences in beneficiary populations between Medicare and CHAMPUS, savings under Medicare might not accrue to CHAMPUS. Thus, while a State may be initially exempt from the CHAMPUS system, we will continue to monitor reimbursement levels in the state to ensure that they do not exceed those under the CHAMPUS DRG-based payment system. If they do exceed that level, we will work with the State to resolve the problem. However, if a satisfactory solution cannot be found, OCHAMPUS will terminate the exemption after due notice has been given to the state.

At this time, it appears that at least one State, Maryland, seeks such an exemption. Such a request will be considered based on these criteria and will likely be approved.

b. Services exempt from the CHAMPUS DRG-based payment system (§ 199.14(a)(1)(ii)(C). In this section we provided a list of specific services which, even if provided in a hospital subject to the CHAMPUS DRG-based payment system, are exempt from the DRG system.

Comment—There is no distinction between the extremely low birth weight patient and those with a higher birth weight but who are still classified as premature. The neonatal DRGs do not have the ability to accommodate the wide variations in case mix severity.

Response-We believe that any significant classification problem with the neonate DRGs occurs primarily at Children's Hospitals which we are exempting from the CHAMPUS DRGbased payment system at this time. Neonates at other hospitals will be included in the system. We have examined the CHAMPUS data and found that the impact of neonate DRGs on hospitals other than children's hospitals is no greater, and maybe even less, than that of other DRGs. For large teaching hospitals, which have only ten percent of the CHAMPUS neonate cases outside children's hospitals, the impact is not significantly greater than for other DRGs. We will pay particular attention to the neonate issue as we analyze CHAMPUS experience under this

Comment—Medicare intends to implement alcohol/drug abuse DRGs and stop exempting such hospitals and units. Will OCHAMPUS do likewise?

Response-We are aware of Medicare's proposed change. However, we believe it would be premature for CHAMPUS to also implement these DRGs at this time because unlike other diagnostic categories we cannot be certain that the nature of alcohol and drug abuse treatment, and its classification, is comparable for both the CHAMPUS and Medicare beneficiary groups. Over the next few months we intend to examine the Medicare changes and evaluate them in terms of CHAMPUS beneficiaries and determine whether it would be appropriate to adopt the Medicare classification changes for the CHAMPUS population or develop a CHAMPUS-specific DRG approach.

Comment—The proposed rule implies that only hospitals may bill for services of hospital-based physicians, but it should depend on the financial arrangement between the physician and the hospital.

Response—A hospital is required to bill for services of hospital-based physicians if the physician is employed by or under contract to the hospital. This requires that the hospital pay the physician. However, if the hospital-based physician merely has an agreement with the hospital to provide services with no requirement that the hospital reimburse the physician, the physician may bill CHAMPUS for his or

her services. We will reword this section to ensure that this is clear.

Comment-The proposed rule apparently includes nurse anesthesia services within the DRG amount. This is a significant departure from the way they are reimbursed under Medicare. The rule should be amended to provide for payment for nurse anesthesia services outside of the DRG system.

Response-When we prepared the proposed rule we were unaware that Medicare had changed from its initial intention to include nurse anesthetists in the DRG payment. We will include a new section which exempts nurse anesthetists from our DRG system and permits hospitals to bill separately for their services. We are aware that Medicare intends to reimburse them as a cost passthrough until October 1989. but CHAMPUS has no comparable costreporting system. Therefore, we are allowing hospitals to bill separately which conforms to the procedures Medicare will use beginning in 1989.

Comment-Bone marrow transplantation and AIDs should be exempt from the CHAMPUS DRG-based

payment system.

Response-Bone marrow transplantation has been an accepted medical practice for sufficient time so that procedures and charges are stablized. Therefore, DRG-based payment amounts accurately reflect average costs. Most admissions of AIDs patients are not for treatment of AIDs. itself, but rather such patients are generally admitted for a complication of AIDs which would be identifiable in our DRG system. Thus, for the present time the DRG system appears able to deal fairly and appropriately with medical care for AIDs and AIDS-related conditions. However, we will continue to monitor developments in this regard and will be ready to make revisions in the future, if appropriate.

c. Hospitals subject to the CHAMPUS DRG-based payment system (§ 199.14(a)(1)(ii)(D)). All hospitals within the fifty states, the District of Columbia, and Puerto Rico which are certified to provide services to CHAMPUS beneficiaries are subject to the DRG-based payment system except for certain types of hospitals or units

which are identified.

Comment-Children's hospitals were specifically excluded from the Medicare DRG-based system due to the inadequacies of DRGs for classifying children's conditions. The DRG classification system does not distinguish the more complex and resource intensive children's conditions which are treated by children's and/or large teaching hospitals. Therefore,

including them would seriously underfund costs for treating these children. They should be exempt.

Response—We have examined CHAMPUS data relating to the charges of children's hospitals, large teaching hospitals and other short-term acutecare hospitals. Our analysis showed that when charges are adjusted for wage, teaching activity, and case mix differences, the average charge per pediatric case for children's hospitals is significantly higher than the national average. Large teaching hospitals and other acute-care hospitals' average charges, however, did not significantly vary from the national average charge per pediatric case. Therefore, we will exempt chidren's hospitals from the CHAMPUS DRG-based payment system because of their unique circumstances.

In order to be exempt, the hospital must be exempt under the Medicare PPS, or if it is not a Medicare provider, it must meet the same criteria required for exemption under the Medicare PPS. However, we believe the CHAMPUS DRG-based payment system can be modified to accommodate children's hospitals, and we intend to review this area so that children's hospitals can be incorporated into our DRG system in the future. We will also welcome any input from interested organizations in this

regard.

Comment-How will exempt facilities be reimbursed?

Response-Hospitals or units which are exempt from the CHAMPUS DRGbased payment system will be reimbursed just as they have been in the past—that is, based on their billed charges.

IV. Determination of Payment Amounts (§ 199.14(a)(1)(iii))

The actual payment for an individual claim under the CHAMPUS DRG-based payment system is calculated by multiplying the adjusted standardized amount by a weighting factor specific to each DRG. The adjusted standardized amounts and the DRG weights shall be calculated from a database of CHAMPUS claims covering at least twelve (12) months.

We used the following procedures to develop the database. CHAMPUS used the same database to calculate both the DRG weights and the adjusted standardized amounts (ASA). The data consisted of all CHAMPUS inpatient claims processed during a 12-month period from July 1, 1986, through June 30, 1987. The data are in UB-82 record format submitted by CHAMPUS fiscal intermediaries.

The database includes only those services and hospitals that will be

subject to DRG reimbursement. Moreover, data errors were removed from the database.

Comment-It is necessary to have the final weights and rates in order to evaluate the impact of the system.

Response-In order to ensure that the final weights and rates are based on the most recent possible data, we are using a database covering July 1, 1986, through June 30, 1987. As a result, the final weights and rates were not available when the proposed rule was published. However, we did calculate preliminary weights and rates from an earlier and more abbreviated database, and we included the resultant adjusted standardized amount and a sample of weights in the proposed rule. This was certainly sufficient to give interested parties a good understanding of the impact of the proposed rule on actual payment amounts, providing additional perspective to aid in consideration of the methodology described in the rule. We also provided a complete list of the preliminary weights to anyone who requested it. The final weights and rates are attached to this final rule for information (Tables 1 and 2). Subsequent changes to the weights and rates will be published as a notice in the Federal Register.

Comment-Several commenters questioned if the data used in the database are reliable.

Response-In our view, there are three components of data reliability: The technical adequacy of the sample, the accuracy of diagnosis and procedure coding, and the reliability of the billed charge amounts on the bill.

We think that the database is technically adequate, because it consists of all CHAMPUS claims processed from July 1, 1986, through June 30, 1987.

The database used for the final weight and adjusted standardized amount (ASA) calculations also has been edited for duplicate records, total charge errors, interim bills, combined bills for mother and newborn services, and admission and discharge date errors. Moreover, all records without valid diagnoses and procedure codes were eliminated from the database. In short, the final database represents a "clean" set of records for the most recently available full year period of CHAMPUS claims. We feel that a dataset of this type will result in reliable and accurate weights and ASA.

We think that the diagnosis and procedure codes are accurate, because they are completed by medical record tehnicians that conduct the same coding for Medicare discharges.

Finally, we are confident that the amount recorded on the bill is the actual billed charges, because the data reflect amounts paid to hospitals, including claims adjustments.

Comment-The proposed October 1 effective date does not permit CHAMPUS to develop a comprehensive database for use in the calculation of weights and the adjusted standardized amount (ASA). Moreover, there isn't sufficient time between the end of the database period and the effective date to calculate the weights and rates.

Response-OCHAMPUS has been working on development of a DRGbased payment system since early 1984. A significant part of this effort has been to develop a comprehensive and accurate database from CHAMPUS claims which could be used to calculate the weights and ASA. We have also previously developed the programming necessary to calculate the weights and ASA. Therefore, the October 1 effective date will have no detrimental effect on the comprehensiveness of the database nor on the ability to timely calculate the weights and ASA

Comment-Since CHAMPUS is ending the database with June 30, the database will not contain adequate data

on long lengths-of-stay.

Response-The database is comprised of claims processed during the subject period and includes final bills. While long stays which are not completed as of June 30 will not be included in the database, any stays which began prior to July 1, 1986, but finished during the database period will be included. There is no reason to believe these will not be comparable. Thus, long stays will be adequately represented in the database.

A. DRG Weighting Factors

1. Calculation of DRG Weights (§ 199.14(a)(1)(iii)(A))

The CHAMPUS DRG weights will be discharge-weighted. Specifically, the denominator used to calculate each weight represents the national average charge per discharge for the average patient. If there are any DRGs which have fewer than ten occurrences in the database, we will use the Medicare weight until we are able to develop a weight based on CHAMPUS data.

Comment-Use of a national average charge per discharge for the weighting denominator, rather than working on a regional or per hospital basis, will have

an adverse effect.

Response-In response to the inequities resulting from other weighting methodologies, Congress required Medicare to adopt the dischargeweighted weights and rates in FY 1988

(section 1886(d)(3)(A) of the Social Security Act, as amended by section 9302(c) of Pub. L. 99-509). Similarly, CHAMPUS will discharge weight its weights and rates.

Comment—Several commenters questioned whether CHAMPUS would be able to calculate accurate weights for all DRGs, or if some DRGs would have insufficient data from which to develop

a weight.

Response-We recognize this as a potential problem. In the proposed rule we proposed using the Medicare weight in those cases where there were no occurrences of a DRG in the database, and in the preliminary database this occurred only twice. However, we realize that a weight for a DRG with only a very few occurrences could be skewed. Therefore, in the calculation of the final weights we have elected to use the Medicare weight for any DRG with fewer than ten occurrences in the database. We believe this action is justified. First, it will ensure that all DRG weights are calculated from an adequate database. Second, our analysis of CHAMPUS weights compared to Medicare weights indicates that on average they are very similar. Third, we expect very few, if any, DRGs to be affected by this.

B. Calculation of the Adjusted Standardized Amount (§ 199.14(a)(1)(iii)(C))

The adjusted standardized amount (ASA) represents the adjusted average operating cost for treating all CHAMPUS beneficiaries in all DRGs during the database period. The ASA does not include any regional or hospital-specific operating cost elements, nor does it contain an urban/ rural distinction.

Comment-Unlike the Medicare PPS, CHAMPUS payment amounts will not be adjusted for differences in prevailing wage levels. This is inappropriate. Wages differ significantly across Metropolitan Statistical Areas (MSAs) and states, reflecting a variety of factors beyond the control of the individual

hospital.

Response—This was one of the most frequent comments. We believe that a DRG-based payment system, whose central premise is payment based on averages, should contain few, if any, adjustments to the payment amounts. Nevertheless, we recognize that wages, although by no means completely beyond the control of hospitals, constitute a large part of hospital costs and vary considerably from area to area. The final rule, therefore, in a significant change from the proposed rule, provides for use of the Medicare

area wage indexes in the CHAMPUS DRG-based payment system.

Medicare finds that after adjusting payments for differences in area wage levels, significant differences in payments still exist between urban and rural hospitals. Our analysis of CHAMPUS claims shows that after adjusting CHAMPUS DRG payments for area wages, the gap between urban and rural payments is substantially narrowed by still warrants an urban/ rural differentiation in the payment system. We have, therefore, included the use of separate urban and rural adjusted standardized amounts in the final rule.

1. Apply the Cost to Charge Ratio (§ 199.14(a)(1)(iii)(C)(1))

Each charge used in the calculation of the ASA is to be reduced to a representative cost by using the Medicare cost to charge ratio which was published in the Federal Register on September 3, 1986 (p. 31523).

Comment-CHAMPUS patients will probably have a different utilization pattern than Medicare patients. Therefore, the ratio is not applicable to

CHAMPUS.

Response-CHAMPUS patients have a different distribution of diagnoses than Medicare patients. CHAMPUS patients, however, due to the fact that they are a younger and healthier population than Medicare patients, require less resource intensity for the same services provided to Medicare patients. For example, hospitals generally post the same room charge for all patients, some of the services provided in this charge include general nursing care and supervision unrelated to specific procedures. In this instance, CHAMPUS patients would require less general nursing care than a more elderly Medicare patient. Therefore, the actual cost or labor intensity required per CHAMPUS patient would be lower than that required of a Medicare patient.

The Medicare cost-to-charge ratio has been derived from extensive research and analysis of actual hospital cost reports. Additionally, diagnosis-specific cost for the CHAMPUS population are not available. We think the Medicare cost-to-charge ratio is the best measure of the relation of hospital costs to charges that is currently available.

2. Increase for Bad Debts (§ 199.14(a)(1)(iii)(C)(2))

The base standardized amount shall be increased by .01 in order to reimburse hospitals for bad debt expenses attributable to CHAMPUS beneficiaries.

Comment-Bad debt on required copayments is not uniformly distributed. some hospitals have higher bad debt levels which should be recognized by CHAMPUS. CHAMPUS should pay actual costs or collect cost-shares themselves.

Response—In the past, CHAMPUS has not reimbursed hospitals for bad debt on beneficiary cost-sharing and deductibles. We think that the inclusion of the same add-on for bad debt to every DRG payment is generous given our past policy. Under this policy, hospitals will be paid prospectively for bad debt expenses for every CHAMPUS discharge through a 1 percentage point add-on to the cost-to-charge ratio.

Moreover, since the large majority of our retiree beneficiaries have either other insurance which is primary or supplemental insurance, we are confident that the allowance is fair and reasonable. Collection of beneficiary cost-shares and deductibles has traditionally been a hospital function and is done for nearly all third-party payers.

3. Preliminary Teaching Standardized Amounts (§ 199.14(a)(1)(iii)(C)(5))

A separate standardized amount shall be calculated for each teaching hospital to reimburse for indirect medical education expenses. This will be done by using a hospital-specific indirect medical education factor calculated in accordance with Medicare procedures.

Although we received no written comments on this, we believe a change from our proposed rule is warranted. Medicare has changed its indirect medical education formula to account for an adjustment for disproportionate share hospitals. This adjustment is scheduled to end September 30, 1989. Since CHAMPUS has no disproportionate share provision, we believe immediate use of the following revised formula is proper.

 $1.5 \times \left[\left\{ 1.0 + \underbrace{number\ of\ interns + residents}_{number\ of\ beds} \right\} .5795 - 1.0 \right]$

Medicare has proposed that Congress reduce the indirect medical education factor in recognition of the fact that the current formula significantly overcompensates hospitals for these costs. CHAMPUS is in agreement with the Medicare proposal, but in order to remain consisent with the current Medicare program, CHAMPUS will use the above formula at this time. Should Congress adopt the Medicare proposal, CHAMPUS will revise its formula accordingly.

4. Updating the Adjusted Standardized Amount (§ 199.14(a)(1)(iii)(C)(7))

Beginning in FY 1989, the ASA will be updated by the Medicare annual update factor, unless the adjusted standardized amount is recalculated.

Comment—It is inappropriate to use the Medicare update factor. CHAMPUS should use the market basket rate of increase.

Response—We think it is important that the government promulgate a uniform DRG update factor for both DRG systems. Medicare and CHAMPUS. Congress establishes this factor each year after considering input from PROPAC, the health care industry, and HCFA. We will comply with the Congressionally-approved update factor.

C. Adjustments to the DRG-Based Payment Amounts (§ 199.14(a)(1)(iv))

Any hospital subject to the CHAMPUS DRG-based payment system can be reimbursed for allowed capital and direct medical education costs upon request. Payment for these costs will be made annually based on the ratio of CHAMPUS inpatient days to total inpatient days.

We have revised this provision in the final rule. In order to conform to the

current statutory requirements for Medicare, the calculated payment for capital costs will be reduced by 7% for FY 1988. If Medicare changes this reduction percentage in the future, CHAMPUS reserves the right to conform to the Medicare change.

Comment—Payment of capital and direct medical education costs annually is inequitable and will hurt hospitals' cash flow, since they must service their debt and pay operating bills timely.

Response—The payments for these items will be sufficiently small, particularly relative to total hospital revenues, that the administrative cost of more frequent payments is not justified. Moreover, CHAMPUS does not have hospital cost information with which to formulate accurate estimates of interim payments for capital and direct medical education.

 Information Necessary for Payment of Capital and Direct Medical Education Costs (§ 199.14(a)(1)(iv)(C))

In order to be reimbursed for allowed capital and direct medical education costs, a hospital must submit a report of its incurred costs to the CHAMPUS fiscal intermediary. The report must cover the same period as the hospital's Medicare cost-reporting period and must be submitted within three months of the end of that period.

Comment—The CHAMPUS DRGbased payment system adds additional reporting requirements. The last thing hospitals need is a separate data gathering-monitoring-reporting system for CHAMPUS beneficiaries.

Response—Without this report there is no way for CHAMPUS to determine what capital and direct medical education costs a hospital has incurred for CHAMPUS beneficiaries. The report

itself is extremely simple nd, in addition to readily available CHAMPUS demographic information, requires only a few other items which are all available from the hospital's Medicare cost report. The CHAMPUS method for reimbursing hospitals for capital and direct medical education results in equitable payments based on hospital-specific CHAMPUS information. Moreover, the reporting requirements of this method are minimal when compared to other capital payment mechanisms such as a cost-finding method.

Comment—All data reported to the FI must agree with the Medicare cost report, and the provider must report any changes. Presently, the Medicare intermediaries supply HCFA with computer tapes/diskettes of all filed cost-reports. Why not have OCHAMPUS and HCFA exchange data?

Response—This would be an enormous administrative task to identify a few changes to the capital and direct medical education payments CHAMPUS makes. In addition, we believe it is the hospital's responsibility to report accurate information to OCHAMPUS.

Comment—CHAMPUS should allow hospitals to use either the Medicare cost-finding method or the aggregate ratio of CHAMPUS PPS-related charges to total charges. A flat per diem is not sensitive to differences in the percentage of operating costs from one ancillary service to another.

Response—The proposed per diem capital payment policy provides reasonable payments for capital costs consistent with the current Medicare per diem payment methodology and results in minimal reporting requirements for hospitals. A cost-finding method similar to Medicare's cost reporting mechanism

is not in hospitals' interest, because it will burden them with extensive reporting requirements for CHAMPUS patients, a payer that generally accounts for a small percentage of each hospital's total revenues. The second alternative, capital payments based on the ratio of CHAMPUS charges to total charges is inadequate. If implemented, this policy would not be consistent with Medicare's current per diem calculation of capital costs. We recognize the fact that Medicare is proposing a revision to their current capital payment policy. CHAMPUS supports this proposal, but due to its reporting requirements on hospitals, it would be administratively infeasible for CHAMPUS to implement at this time. When the Medicare proposal is finally implemented in the Medicare system, CHAMPUS will consider it for inclusion in the CHAMPUS DRG-based payment system.

2. Outliers (§ 199.14(a)(1)(iii)(D))

CHAMPUS will adjust DRG-based payments for atypical cases. These outliers are those cases that have either an unusually short length-of-stay or extremely long length-of-stay or that involve extraordinarily high costs when compared to most discharges classified in the same DRG.

Comment-There is no mention of a

target outlier pool.

Response—The CHAMPUS DRGbased payment system does not have an outlier pool. Under the Medicare PPS, an estimated percentage of total DRGbased payments is set aside as an outlier pool, thus actually reducing the non-outlier DRG-based payments. Under the CHAMPUS DRG-based payment system, outlier payments are made over and above the basic DRG payments.

Comment—Medicare is changing their outlier policy to recognize the most expensive cases. Will CHAMPUS make

similar changes?

Response—We do not intend to adopt the proposed changes until we can fully assess their impact on CHAMPUS claims and whether they produce more equitable payments.

Comment—Under the CHAMPUS DRG system, providers are at unlimited

risk for extremely ill patients.

Response—Providers will not be at unlimited risk, because our system includes payments for outliers, and we believe the outlier payments are reasonable.

Comment—Will the outlier thresholds be revised to coincide with the Medicare

outlier thresholds?

Response—There is no intent that the thresholds be the same. While our methodology for determining the

thresholds is the same, the actual thresholds will be derived from the CHAMPUS database and will be different. Our intent is that the thresholds reflect the service patterns for CHAMPUS beneficiaries.

a. Short-stay outliers
(§ 199.14(a)(1)(iii)(D)(1)(i)). Any
discharge with a length-of-stay (LOS)
less than 1.94 standard deviations from
the DRG's geometric LOS shall be
classified as a short-stay outlier. Shortstay outliers will be reimbursed at 200
percent of the per diem rate for the DRG
for each covered day of the hospital
stay.

Comment—Several commenters objected to our inclusion of short-stay outliers in the CHAMPUS DRG-based

payment system.

Response—Short-stay outliers are fully justified. Just as an exceptionally long length-of-stay should not be considered typical for a given DRG and deserves additional payment, an exceptionally short stay should also not be considered typical and payment should be reduced. Nevertheless, we recognize that the initial days of a hospital stay are generally more expensive, so we have set the short-stay reimbursement at 200 percent of the per diem.

Comment—Many patients will have lengths-of-stay just over the short-stay outlier cutoff. This will make the process of medical review extremely expensive for both CHAMPUS and the provider.

Response—Since DRG-based payment is based on averages, we expect many stays to be shorter than the average. The short-stay outlier policy is intended to isolate those cases with a length-of-stay which is so different from the norm (in this case extraordinarily short) that it is not representative of cases within that DRG. However, the fact that cases exceed the short-stay cutoff but still are less than average will not affect the amount or extent of review to be performed.

Comment—It was indicated an average payment amount was used because some cases will cost more than others, but it will even out in total. Short-stay outliers eliminate that

possibility.

Response—This is not correct. The short-stay outlier policy only eliminates those cases with such unusually short lengths-of-stay, that they are not representative of cases within the DRG. However, many cases will exceed the short-stay cutoff and still be less than the average length-of-stay. It is these cases which will allow the averaging effect. The short-stay policy is simply a counterbalance to the long-stay policy which is well accepted.

Comment—The short-stay outlier policy could result in significant financial losses in cases where a patient dies soon after admission.

Response—We recognize the fact that the initial days of a hospital stay are relatively more costly than the final days of a stay. Therefore, we will pay hospitals 200 percent of a DRG-specific per diem for short stay outlier cases. We think that this level of payment is reasonable.

V. Charges to Beneficiaries

A. Inpatient Cost-Sharing

 Services Subject to the CHAMPUS DRG-Based Payment System (§ 199.4(f)(3)(ii)(A))

Under the proposed rule, for beneficiaries other than dependents of active-duty members the cost-share would have been a per diem amount for each day of the hospital stay, except that the day of discharge would not be counted. The per diem amount would be calculated so that total cost-sharing amounts for these beneficiaries is equivalent to 25 percent of the CHAMPUS-determined allowable costs for hospital services.

Within the Department of Defense there has been some concern that the proposed cost-sharing policy could result in some beneficiaries paying more cost-share than they would pay under current procedures. We therefore examined data which projected cost-shares under both the current and proposed procedures, and we found that approximately 30% of these beneficiaries would have greater cost-shares, although in most cases the actual increase was quite small. Even so, we found this to be an unacceptable consequence for our beneficiaries.

In order to correct this situation, the final rule modifies the cost-sharing provisions for these beneficiaries. The cost-share will be calculated at the lesser of the per diem-based amount or 25 percent of the hospital's billed charges. In this way no beneficiary will be responsible for a larger cost-share than currently required, and most will have smaller cost-shares. As a result of this change, the per diem amount has been increased slightly so that total cost-sharing amounts still equal 25% of the CHAMPUS-determined allowable amounts as required by the CHAMPUS statute.

This change in policy from the proposed rule does not in any way reduce the overall substantial cost-savings for beneficiaries as a whole. Non-active duty dependent beneficiaries will pay significantly less than they do

now. The change in policy means that none of these beneficiaries will pay any more than they do now; most will pay less; and that about half of the time they will save over \$100 per hospital stay.

Comment—What is the beneficiary cost-share if the patient dies on the day of admission?

Response—In any case in which the length-of-stay is less than 24 hours, but the stay qualifies as an inpatient stay under CHAMPUS rules, the beneficiary cost-share is to be calculated based on a one-day stay.

Comment—It will be cumbersome to calculate beneficiary cost-shares when the actual charges are less than the DRG-based payment.

Response—We have been very careful to ensure that the cost-sharing methodology would be easy for hospitals to calculate at the time of discharge. Since both length-of-stay and actual charges are known at discharge, the hospital should have no trouble calculating the beneficiary's cost-share.

B. Hospital Days Beyond that Deemed Medically Necessary (§ 199.4(f)(6))

Under current CHAMPUS procedures, as required by law, no CHAMPUS payments may be made for care provided which is not medically necessary. Although it does not occur frequently, application of this rule may result in beneficiaries being responsible for payments they hoped CHAMPUS would cover. Some media reports after publication of the proposed rule created confusion regarding application of this requirement under the DRG system.

Under the DRG-based payment system, this requirement is unchanged. In the usual case, the DRG amount will be considered full payment for all days of care. However, in the unusual cases in which the length of stay exceeds the long-stay outlier standard, additional payments will generally be made to the hospital. In these cases, it is possible that applying the requirement of medically necessary care could result in some or all of the additional payment being disallowed. In addition, as in the current program, an entire hospital admission could be determined not medically necessary, resulting in the claim being disallowed.

These possibilities, however, do not represent any change from currently required procedures. Moreover, implementation of the DRG-based payment system is not expected to result in any increase in the frequency of disallowed claims based on a lack of medical necessity.

VI. Quality of Care Review (§ 199.14(a)(1)(v))

Implementation of the Medicare DRG system generated concerns about the quality of care. In response to these concerns Congress established a Peer Review Organization (PRO) system to determine adequacy and appropriateness of care. Congressional hearings continued to be held that focused on complaints of premature discharge. Medicare revised its PRO system to concentrate further on quality of care issues. As a result of these efforts, the state of the art of process. structure, and outcome measures of the quality of care has evolved to the point where both nationally and locally developed peer review systems are widely recognized as effectively monitoring the quality of care provided.

In connection with implementing the CHAMPUS DRG system, we will piggyback on the established Federal system for monitoring the quality and appropriateness of civilian inpatient care for its beneficiaries.

CHAMPUS will implement a quality of care review program which will assure the appropriateness of care provided to beneficiaries. Every case reimbursed under the DRG system will be subject to generic quality screen reviews, admission and discharge reviews, and DRG validation. One of the objectives of these multiple reviews is to guard against episodes of premature discharge or inappropriate admission. A peer review system will use criteria which has been developed on both national and local levels to determine the adequacy and appropriateness of care. This system, which is modeled on the Medicare PRO system, will be specific to the CHAMPUS population.

Comment—The current CHAMPUS system for reviewing psychiatric and substance abuse treatment works well and ensures adequate and qualified review. Therefore, psychiatric and substance abuse services should be excluded from PRO review.

Response—The admission and quality review system set forth in this rule applies only to those services reimbursed under the CHAMPUS DRG-based payment system. Therefore, psychiatric and substance abuse services, which are exempt from DRG-based payment, will be excluded from the review system.

Comment—What are the procedures to be followed in conducting the admission and quality review?

Response—There will be two phases of review under the CHAMPUS DRG-based payment system. The permanent review system is being coordinated with

the Health Care Financing Administration. HCFA is currently in the process of recomputing its PRO contracts and the new contracts will be phased in between July and December 1988. Quality assurance reviews of CHAMPUS claims and services may be included in future PRO contracts. In the interim, CHAMPUS will conduct quality of care reviews under a separate contract. The admission and quality review section of this rule provides the basis for these review functions. Information regarding the detailed procedures will be found in the RFPs and resulting contracts for both the interim an proposed permanent review systems.

A. Areas of Review

1. Admissions (§ 199.14(a)(1)(v)(C)(1))

This section sets forth the areas which are to be reviewed to determine whether inpatient care is medically necessary and whether services are delivered in the most appropriate setting.

Comment—All transfers of CHAMPUS beneficiaries from a hospital or hospital unit subject to the CHAMPUS DRG-based payment system to another hospital or hospital unit are to be reviewed. Does this include transfers from a hospital or unit subject to the DRG system to a hospital or unit exempt from the system?

Response—We have clarified in this final rule that these cases are to be classified as discharges. This particular section applies only to those actions which are classified as transfers, so a transfer to a hospital or unit exempt from the DRG system would not be reviewed under this provision.

Comment—All CHAMPUS admissions to a hospital or unit subject to the DRG system which occur within seven calendar days of discharge from a hospital or unit subject to the DRG system are to be reviewed. Will the time periods involved in this provision be changed to coincide with Medicare's review requirement?

Response—To the extent possible, we want hospitals and PROs to be subject to only a single set of review procedures. Therefore, we will use the same time periods.

Comment—If the number of unnecessary CHAMPUS admissions for a hospital is more than 2.5 percent of the review sample or three cases (whichever is greater) for any quarter, all CHAMPUS admissions for that hospital must be reviewed during the following quarter. Will this threshold for intensified review be changed to coincide with Medicare? Can the review

organization focus on identified subsets rather than intensify an entire provider?

Response—The requirements will

duplicate Medicare's.

Comment—Prepayment review is required for all CHAMPUS admissions in any DRGs which have been specifically identified by OCHAMPUS. Has OCHAMPUS identified any such DRGs?

Response—None have been identified to date. The selection of the DRGs for pre-admission or pre-procedure review will be based on data which targets known or suspected high risk topics.

2. Admission Pattern Monitoring (§ 199.14(a)(1)(v)(C)(2))

In order to ensure that discharges are appropriate, admissions for those hospitals identified as having significant increases in quarterly discharges shall be reviewed.

Comment—Medicare has determined that admission pattern monitoring was nonproductive and has eliminated it. Will CHAMPUS?

Response—As for other areas of review, we will duplicate Medicare in this respect.

3. Procedure Review (§ 199.14(a)(1)(v)(C)(5))

All claims for procedures identified by OCHAMPUS as subject to a pattern of abuse shall be reviewed.

Comment—Has OCHAMPUS identified any procedures subject to this review?

Response—Not yet. This type of review will not be implemented until the permanent review system has begun.

B. Fiscal Intermediary Actions as a Result of Review (§ 199.14(a)(1)(v)(D))

CHAMPUS intends for the FI, not the PRO, to institute corrective actions by the hospital. Therefore, this section enumerates those actions which a fiscal intermediary may take if the PRO determines that a hospital has misrepresented admission, discharge, or billing information, or has taken an action that results in the unnecessary admission of an individual entitled to benefits, unnecessary multiple admissions of an individual, or other inappropriate medical or other practices. This finding may be with regard to an individual claim or a pattern of inappropriate practices.

inappropriate practices.

Comment—What procedures have been developed for this process—how is the fiscal intermediary to be notified,

Response—As for other detailed information on these procedures, it will be found in the RFPs and resulting contracts for both the interim and

permanent review systems.

OCHAMPUS also will issue guidance in the future on the review systems which will provide additional information.

VII. Summary of Regulations Changes

For the convenience of the reader, we are summarizing the changes we are making to the proposed rule as a result of public comments. The reader is referred to the detailed discussions above for a complete explanation of the rationale for these changes.

A. Urban/Rural Differentiation

There will be separate adjusted amounts (ASAs) calculated for urban and for rural areas. The same urban/rural designations used in the Medicare PPS will be used in the CHAMPUS DRG payment system.

B. Wage Index

Each adjusted standardized amount (ASA) which will be divided into labor and nonlabor portions. The wage index adjustments used by Medicare will be applied to the labor portions of each ASA. This amount will then be added to the nonlabor portion, and the sum will be multiplied by the DRG weight to arrive at the DRG-based payment amount.

C. State Waivers

Any state which has been exempted from the Medicare PPS can request an exemption from the CHAMPUS system.

D. Children's Hospitals

Any children's hospital which is exempt from the Medicare PPS will be exempt from the CHAMPUS system.

E. Cost-Shares

The cost-share for beneficiaries other than dependents of active duty members will be the lesser of (a) the amount based on the per diem as described in the proposed rule, or (b) 25% of the billed charge, but it can never exceed the DRG-based amount.

F. Grouper Program

The CHAMPUS system will use the most recently available Medicare Grouper program.

G. Nurse Anesthetists

Hospitals will be allowed to bill separately for nurse anesthetists' services just as they may for hospitalbased physicians.

H. Capital and Direct Medical Education Payments

The capital payment for FY 1988 is to be reduced by 7% in accordance with Medicare procedures.

I. Discharges/Transfers

It is to be considered a discharge if the patient is transferred from the care of a hospital included under the CHAMPUS DRG-based payment system to a hospital or unit that is excluded from the DRG system, except for transfers to uniformed services treatment facilities.

VIII. Summary of Differences from the Medicare PPS

Although the CHAMPUS DRG-based payment system is modeled on the Medicare PPS, there are several differences. Below we have summarized the significant differences which are not addressed in the summary of public comments.

A. Services Subject to the CHAMPUS DRG-Based Payment System

The CHAMPUS DRG-based payment system exempts some services which are included in the Medicare PPS. They are: Psychiatric services in short-term hospitals; heart transplantation services; and liver transplantation services. We have exempted psychiatric services, because research has shown that psychiatric DRGs may not be a reliable measure of cost variability for psychiatric cases under CHAMPUS. We have also exempted heart and liver transplantation services for much the same reason—that is, we are concerned that the DRG-based amounts also may not be a reliable measure of their cost variability, since they occur infrequently and involve significant costs.

B. Hospitals Subject to the CHAMPUS DRG-Based Payment System

CHAMPUS has exempted certain types of hospitals from our DRG system which are subject to the Medicare PPS. Sole community hospitals are paid under special provisions under the Medicare PPS based in part on hospitalspecific costs which are unavailable to CHAMPUS. Therefore, we have exempted them. Christian Science Sanitoriums are paid a predetermined fixed amount per discharge under Medicare. However, since they involve such a small number of providers and CHAMPUS claims and since some may qualify under the long-term hospital exemption anyway, we have elected to exempt all of them.

C. Updating DRG Weights

Medicare is required to update their weights annually. We plan to recalculate CHAMPUS weights annually based on a charge sample from the most recent period under CHAMPUS prospective payment. Notice of the

revised weights will be published in the Federal Register.

D. Inclusion of Puerto Rico

Although Puerto Rico is included in the Medicare PPS beginning in October 1987, hospitals there will be reimbursed using a blend of the national rate and the Puerto Rico discharge-weighted urban or rural standardized rate. The CHAMPUS DRG-based payment system makes no distinction for Puerto Rican hospitals.

E. Capital and Direct Medical Education Payments

These items are reimbursed as a cost passthrough under the Medicare PPS. CHAMPUS will pay for these items upon written request. CHAMPUS has no cost-reporting mechanism, but the report of these costs which is submitted to the CHAMPUS fiscal intermediary must correspond to the Medicare cost-report. We support the current Medicare proposals that revise the payment methodologies for capital and direct medical education. CHAMPUS' implementation of these proposals at this time would be administratively infeasible. However, once Medicare adopts the new methodologies. CHAMPUS will consider them for inclusion in the CHAMPUS system.

F. Bad Debt Adjustment

The adjusted standardized amount used in the CHAMPUS DRG-based payment system contains a factor to reimburse hospitals for CHAMPUS' share of their bad debts. Under the Medicare PPS, bad debts are reimbursed as a cost passthrough.

IX. Impact Analysis

A. Executive Order 12291 and the Regulatory Flexibility Act

Executive Order 12291 requires that a regulatory impact analysis be performed on any major rule. A "major rule" is defined as one which would:

Result in annual effect on the national economy of \$100 million or more;

Result in a major increase in costs or prices for consumers, any industries, any government agencies, or any geographic regions; or

Have significant adverse effects on competition, employment, investment, productivity, innovation or on the ability of U.S.-based enterprises to compete with foreign-based enterprises in domestic or import markets.

The Regulatory Flexibility Act requires that each federal agency prepare, and make available for public comment, a regulatory flexibility analysis when the agency issues regulations which would have a

significant impact on a substantial number of small entities. For purposes of the Regulatory Flexibility Act, we consider small entities to include all nonprofit and most for-profit hospitals.

Under both the Executive Order and the Regulatory Flexibility Act, such analyses must, when prepared, examine regulatory alternatives which minimize unnecessary burden or otherwise assure that regulations are cost-effective.

We are treating this final rule as a major rule under Executive Order 12291, since we anticipate that the changed reimbursement procedures required by this final rule will result in annual program savings exceeding \$100 million. The Department of Defense Authorization Act, 1984, which provides the authority for CHAMPUS to use a DRG-based payment system, allows some administrative discretion in the implementation of such a reimbursement system. Therefore, this analysis examines the major features of the system and the rationale for each.

Because of the extensive changes this final rule will cause in our methods for paying for inpatient hospital services, we are providing the following discussion which, when combined with the rest of this preamble, constitutes a regulatory impact analysis and a voluntary regulatory impact/flexibility analysis.

B. The Problem of Increased CHAMPUS

The rapidly rising costs of health care have been the focus of numerous studies and have resulted in many efforts to curb the rise. Most notable of these efforts is the implementation of the Medicare Prospective Payment System (PPS) which was implemented in October 1983. Although the PPS was required to be "budget neutral" during its early years, it has had a significant impact, not only on Medicare, but also on the delivery of health care services to the public as a whole. CHAMPUS has unquestionably benefited from this in certain respects, but nevertheless our costs continue to rise at an unacceptable rate. For example, a comparison of CHAMPUS data for FY 1985 to FY 1983 shows that, while the number of admissions and the average length-ofstay have decreased, the cost per admission has increased 19.4 percent. the cost per inpatient day has increased 26.2 percent and total CHAMPUS expenditures for inpatient care increased 11.1 percent. This trend continued into FY 1986 with total hospital costs increasing 19.0 percent from FY 1985, although admissions during that year also increased by 9

percent. Average length-of-stay remained the same.

We attribute these increases to several factors. The first is inflation, but since inflation in the economy as a whole has slowed considerably, its role in the increases is minor. A second contributing factor is the absence of traditional supply and demand forces operating to curb excessive expenditures, although, like inflation, this has been checked somewhat in recent years, particularly by the Medicare PPS and other similar programs. A third factor which is significant is CHAMPUS' practice of reimbursing hospitals based on their billed charges. This creates no incentive for hospitals to control costs, and, in fact, creates the opposite incentive. This ties into the fourth factor, which is costshifting to billed charge payers such as CHAMPUS from other third-party payers which have placed limitations on payments.

C. A DRG-Based Payment System Represents the Best Resolution of the Problem of Increasing Costs

There can be no doubt that the Medicare PPS has significantly affected the delivery of hospital services in the United States. The CHAMPUS DRGbased payment system closely resembles the Medicare system and will benefit from the same advantages. Of particular importance, it will enable us to set our reimbursement levels to more closely equal hospitals' costs of providing services to our beneficiaries, and it will enable us to avoid the increases in charges resulting from costshifting which results in CHAMPUS subsidizing non-CHAMPUS patients. We fully intend to reimburse hospitals the reasonable costs of providing care to our beneficiaries, but in order to maintain the level of benefits offered by CHAMPUS under increasing budgetary constraints, it is incumbent upon us to implement steps to control our costs.

D. Quantification of Impact

In our initial impact analysis in the proposed amendment of rule we described the impacts we expect the CHAMPUS DRG-based payment system to have on hospitals, beneficiaries, and fiscal intermediaries. We also solicited comments and factual information that would enable us to describe and quantify in greater detail the effects of our DRG system. Although we received numerous comments regarding specific provisions which we have addressed earlier in this preamble, we received no specific information with regard to the economic impact.

Subsequent to publication of the proposed rule we have completed a further analysis of the impact of this change. This analysis resulted in an estimate of \$200 million savings to CHAMPUS for inpatient services provided during FY 1988. However, the effects of DRG-based payments will not actually begin until perhaps three months after implementation of the system because of actual service time, time to submit claims, and the time to process the claims. Therefore, actual CHAMPUS savings during FY 1988 would be approximately \$150 million.

E. Economic Impacts

In this section we will discuss the impact on hospitals, on our beneficiaries, and on CHAMPUS operations.

1. Hospital Impact

Since the Medicare PPS has been in operation for nearly four years, we believe hospitals have adjusted their operations to accommodate it. Therefore, we anticipate few, if any, changes in hospital operations as a result of our implementation of the CHAMPUS DRG-based payment system. Possible operational changes could be in hospitals' billing practices (if they have not already adopted Medicare billing requirements for CHAMPUS), in the need for additional medical records personnel, and perhaps in increased hospital utilization review activities. For the most part, however, these impacts should be negligible, since CHAMPUS beneficiaries constitute such a small part of most hospitals' patient loads. There may be some hospitals which serve a large number of CHAMPUS beneficiaries and relatively few Medicare beneficiaries, and the impact on these hospitals' operations may be greater, but we expect such hospitals to be very few.

Moreover, we believe some of the distinctions used during the Medicare PPS transition period, such as hospitalspecific and regional differentiations, are not necessary for the CHAMPUS DRG-based payment system, because CHAMPUS charges generally represent a small percentage of each hospital's total revenues and because hospitals have already adjusted their operating practices in response to the Medicare PPS. Moreover, the end of Medicare's phase-in period which recognized the regional and hospital-specific distinctions will approximately coincide with the effective date of the CHAMPUS DRG-based payment system. However, as noted earlier in this preamble, we have included an urban/rural differentiation and a wage index

provision in the final rule in order to recognize these differences.

The primary impact of the CHAMPUS DRG-based payment system will be in the immediate reduction of total CHAMPUS payments to hospitals. It will also give us the ability to control increases in costs in the future. Because the CHAMPUS DRG-based payment system is modeled on the Medicare PPS, CHAMPUS payments for our beneficiaries will be more proportionate to Medicare payment for Medicare beneficiaries. In addition, CHAMPUS will no longer pay those amounts which have been shifted to charge payers because of payment limitations imposed by various states and other third-party

The CHAMPUS DRG-based payment system includes a number of provisions and procedures which we believe help to mitigate its impact on hospitals. These include:

a. Use of Medicare cost to charge ratio. The base from which the standardized amounts are calculated is 66 percent of charges. This is the Medicare cost to charge ratio which was published in the Federal Register on September 3, 1986. This ratio excludes capital costs and direct medical education costs. Because hospitals' posted charges to third-party payers are generally consistent, this ratio provides a reasonable estimate of CHAMPUS costs relative to charges. Moreover, it represents those costs which have been identified, through statute and regulation, as reimbursable under the major government program. At the same time, this ratio is derived from claims for Medicare beneficiaries. Since our beneficiaries are considerably younger and generally healthier on average, we believe that an average CHAMPUS beneficiary would use fewer hospital resources than an average Medicare beneficiary classified under the same

b. Bad debts. In order to recognize our share of hospitals' bad debts for CHAMPUS patients (that is, unpaid cost-sharing amounts), we have increased the base amount for the standardized amounts from .66 to .67. This is an increase of about 1.5 percent which is actually more than our share of CHAMPUS bad debts.

c. Use of CHAMPUS-specific weights. We recognize that, because of the differences between our beneficiaries and Medicare's beneficiaries, their relative resource consumption in the various DRGs will be different. In order to ensure that the payment amounts used in the CHAMPUS DRG-based payment system are reasonable for our

beneficiaries, we will calculate DRG weights form CHAMPUS claims data only.

d. Wage index. We recognize that wages, although by no means completely beyond the control of hospitals, constitute a large part of hospital costs and vary considerably from area to area. We have, therefore, provided for use of the Medicare area wage indexes in the CHAMPUS DRGbased payment system. This provision is essentially budget neutral for CHAMPUS, since it will not directly affect the total amounts paid. However, it will redistribute those payments so that, rather than all hospitals being paid the same amount for the same DRG. those hospitals in high wage areas will be paid more than the average and those hospitals in low wage areas will be paid less.

e. Urban/rural differentiation.
CHAMPUS data indicate that significant differences in the impact of DRG-based payments exist between urban and rural hospitals even after adjusting for area wage differences. We have, therefore, provided for development of separate urban and rural ASAs based on claims data specific to each. As for the use of wage indexes, this provision will be hudget neutral

f. Capital and direct medical education costs. In one CHAMPUS study of the largest inpatient CHAMPUS hospitals, we found that, as a percentage of total expenses, capital costs ranged from 2.4 percent to 20.5 percent. In the same study we found that direct medical education costs vary from 0 percent to 6.1 percent of total expenses. We recognize these are expenses which apply to our beneficiaries. Moreover, at present there is no equitable way to reimburse hospitals for these costs on a uniform basis which would not unduly penalize certain hospitals. The CHAMPUS DRGbased payment system, therefore, includes procedures for hospitals to report their total capital and direct medical education costs to CHAMPUS and be reimbursed CHAMPUS' share based on the ration of CHAMPUS inpatient days to total inpatient days.

g. Phychiatric services. Most psychiatric services are exempt from DRGs under Medicare with the exception of psychiatric services provided in some acute care facilities that do not qualify for an exemption. Initially, all psychiatric services are exempt from the CHAMPUS DRG-based payment system, but OCHAMPUS is currently studying whether other prospective payment methodologies are appropriate for the provision of

psychiatric services to CHAMPUS patients.

h. Substance abuse services. Although Medicare has developed a DRG classification system for substance abuse services adequate for its beneficiary population, at this time we think that it is unclear whether this system should be applied to CHAMPUS patients because of the distinct characteristics of the CHAMPUS population. Therefore, we have elected to conduct further study on this issue. In the meantime, we will exempt substance abuse services from the CHAMPUS DRG-based payment system and continue to pay for them according to billed charges.

i. Children's hospitals. Both the comments we received, and CHAMPUS data we have examined, indicate that children's hospitals have higher costs. on average, than other hospitals. As a result, we have exempted children's hospitals from the CHAMPUS DRG-

based payment system.

2. Beneficiary Impact

The CHAMPUS DRG-based payment system will benefit our beneficiaries and the procedures in this final rule contain various provisions to protect them.

a. Cost-sharing amounts. The costsharing provisions under the CHAMPUS DRG-based payment system are structured so that beneficiaries are still responsible on average for the same proportion of allowed costs. There will be no effect on dependents of active duty members in this regard, but on average all other beneficiaries will be required to pay smaller cost-sharing amounts, since the allowed amounts will be reduced.

b. Calculation of cost-shares for beneficiaries other than dependents of active duty members. We conducted a test of DRG-based reimbursement in South Carolina from September 1, 1984, through August 31, 1985. One of the most significant findings of that test was that the calculation of cost-shares for beneficiaries other than dependents of active duty members must be revised. Currently these beneficiaries' cost-share is 25 percent of the allowed amount which is generally nearly equal to the billed amount. During the test we found that under a DRG-based payment system the DRG-based amount sometimes greatly exceeded the hospital's billed charge, resulting in a cost-share equal to, and sometimes exceeding, the billed charge. In order to prevent this inequity, we revised the cost-sharing procedures for these beneficiaries in the proposed rule to require a standard per diem amount for services provided by hospitals subject to

the CHAMPUS DRG-based payment system. This did not entirely correct the problem, however, and some beneficiaries would still be required to pay cost-shares exceeding what would be required under current procedures. Thus, this final rule further modifies the cost-sharing requirements for these beneficiaries. The cost-share will now be the lesser of the per diem-based amount or 25 percent of the hospital's billed charges. In this way no beneficiary's cost-share will increase as a result of DRG-based payment (see Section IV.A. of this preamble).

3. Operational Impact

Fiscal intermediaries will have to make significant changes to their existing claims processing systems in order to implement the CHAMPUS DRGbased payment system. An OCHAMPUS negotiation team has been established to negotiate reimbursement of costs with the fiscal intermediaries.

F. Conclusion

We believe that this final rule meets the objectives of the Regulatory Flexibility Act.

X. Other Required Information

A. Effective Date

The procedures contained in this final rule are to be applicable to all admissions occurring on or after October 1, 1987. In the proposed amendment of rule we stated that implementation of DRG-based reimbursement in Hawaii would be delayed until April 1988. This was based on potential delays in making administrative adjustments for fiscal intermediary activities in Hawaii. These issues have now been resolved and there is no administrative reason for a delay. Thus, we no longer intend to postpone implementation in Hawaii, and DRG-based reimbursement will be implemented there on October 1, 1987.

B. Paperwork Reduction Act

This final rule contains a reporting requirement for capital and direct medical education costs which is subject to the provisions of the Paperwork Reduction Act of 1980 (44 U.S.C. 3507) As required by that Act, OCHAMPUS has requested Office of Management and Budget (OMB) approval of this requirement.

List of Subjects in 32 CFR Part 199

Claims, Handicapped, Health insurance, Military personnel.

PART 199-[AMENDED]

Accordingly, 32 CFR Part 199 is amended as follows:

1. The authority citation for Part 199 continues to read as follows:

Authority: 10 U.S.C. 1079, 1086, 5 U.S.C. 301.

2. Section 199.2(b) is amended by adding the following definitions in alphabetical order:

§ 199.2 Definitions.

CHAMPUS DRG-Based Payment System. A reimbursement system for hospitals which assigns prospectivelydetermined payment levels to each DRG based on the average cost of treating all CHAMPUS patients in a given DRG.

Diagnosis-Related Groups (DRGs). Diagnosis-related groups (DRGs) are a method of dividing hospital patients into clinically coherent groups based on the consumption of resources. Patients are assigned to the groups based on their principal diagnosis (the reason for admission, determined after study). secondary diagnoses, procedures performed, and the patient's age, sex, and discharge status.

3. § 199.4 is amended by revising paragraphs (d)(2), (f)(3)(ii), (f)(4)(ii), (f)(5), (g)(10), and (g)(11) and by adding paragraph (f)(6) to read as follows:

§ 199.4 Basic program benefits.

(d) * * *

(2) Billing practices. To be considered for benefits under this paragraph (d). covered services and supplies must be provided and billed for by an authorized provider as set forth in § 199.6 of this part. Such billing must be itemized fully and described sufficiently, even when CHAMPUS payment is determined under the CHAMPUS DRG-based payment system, so that CHAMPUS can determine whether benefits are authorized by this part. Except for claims subject to the CHAMPUS DRGbased payment system, whenever continuing charges are involved, claims should be submitted to the appropriate CHAMPUS fiscal intermediary at least every 30 days (monthly) either by the beneficiary or sponsor or directly by the provider. For claims subject to the CHAMPUS DRG-based payment system, claims may be submitted only after the beneficiary has been discharged or transferred from the hospital.

(f) * * *

(3) * * *

 (ii) Inpatient cost-sharing. Costsharing amounts for inpatient services shall be as follows:

(A) Services subject to the CHAMPUS DRG-based payment system. The cost-share shall be the lesser of: an amount calculated by multiplying a per diem amount by the total number of days in the hospital stay except the day of discharge; or 25 percent of the hospital's billed charges. The per diem amount shall be calculated so that, in the aggregate, the total costsharing amounts for these beneficiaries is equivalent to 25 percent of the CHAMPUS-determined allowable costs for covered services or supplies provided on an inpatient basis by authorized providers. The per diem amount shall be published annually by OCHAMPUS.

(B) Services exempt from the CHAMPUS DRG-based payment system and services provided by hospitals and parts of hospitals exempt from the CHAMPUS DRG-based payment system and by institutions other than hospitals. The cost-share shall be 25 percent of the CHAMPUS-determined allowable costs or charges for otherwise covered services or supplies provided on an inpatient basis by an authorized provider.

(4) * * *

(ii) Inpatient cost-sharing. Eligible former spouses are responsible for the payment of cost-sharing amounts the same as those required for retirees, dependents of retirees, dependents of deceased active duty members, and dependents of deceased retirees.

(5) Amounts over CHAMPUSdetermined allowable costs or charges. It is the responsibility of the CHAMPUS fiscal intermediary to determine allowable costs for services and supplies provided by hospitals and other institutions and allowable charges for services and supplies provided by physicians, other individual professional providers, and other providers. Such CHAMPUS-determined allowable costs or charges are made in accordance with the provisions of § 199.14. All CHAMPUS benefits, including calculation of the CHAMPUS or beneficiary cost-sharing amounts, are based on such CHAMPUS-determined allowable costs or charges. The effect on the beneficiary when the billed cost or charge is over the CHAMPUSdetermined allowable amount is dependent upon whether or not the applicable claim was submitted on a participating basis on behalf of the

beneficiary or submitted directly by the beneficiary on a nonparticipating basis and on whether the claim is for inpatient hospital services subject to the CHAMPUS DRG-based payment system. This provision applies to all classes of CHAMPUS beneficiaries.

Note.—When the provider "forgives" or "waives" any beneficiary liability, such as amounts applicable to the annual fiscal year deductible for outpatient services or supplies, or the inpatient or outpatient cost-sharing as previously set forth in this section, the CHAMPUS-determined allowable charge or cost allowance (whether payable to the CHAMPUS beneficiary or sponsor, or to a participating provider) shall be reduced by the same amount.

(i) Participating provider. Under CHAMPUS, authorized professional providers and institutional providers other than hospitals have the option of participating on a claim-by-claim basis. Participation is required for inpatient claims only for hospitals which are Medicare-participating providers. Hospitals which are not Medicareparticipating providers but which are subject to the CHAMPUS DRG-based payment system in paragraph (a)(1) of § 199.14 must sign agreements to participate on all CHAMPUS inpatient claims in order to be authorized providers under CHAMPUS. All other hospitals may elect to participate on a claim-by-claim basis. Participating providers must indicate participation by signing the appropriate space on the applicable CHAMPUS claim form and submitting it to the appropriate CHAMPUS fiscal intermediary. In the case of an institution or medical supplier, the claim must be signed by an official having such authority. This signature certifies that the provider has agreed to accept the CHAMPUSdetermined allowable charge or cost as payment in full for the medical services and supplies listed on the specific claim form, and further has agreed to accept the amount paid by CHAMPUS or the CHAMPUS payment combined with the cost-sharing amount paid by or on behalf of the beneficiary as full payment for the covered medical services or supplies. Therefore, when costs or charges are submitted on a participating basis, the patient is not obligated to pay any amounts disallowed as being over the CHAMPUS-determined allowable cost or charge for authorized medical services or supplies.

(ii) Nonparticipating providers.

Nonparticipating providers are those providers who do not agree on the CHAMPUS claim form to participate and thereby do not agree to accept the CHAMPUS-determined allowable costs or charges as the full charge. For

otherwise covered services and supplies provided by such nonparticipating CHAMPUS providers, payment is made directly to the beneficiary or sponsor and the beneficiary is liable under applicable law for any amounts over the CHAMPUS-determined allowable costs or charges. CHAMPUS shall have no responsibility for any amounts over allowable costs or charges as determined by CHAMPUS.

determined by CHAMPUS.

(6) Hospital days beyond that deemed medically necessary. Under the CHAMPUS DRG-based payment system, the DRG amount is considered full payment for any hospital stay up to the long-stay outlier cutoff as described in paragraph (a)(1)(iv)(D)(1)(ii) of § 199.14. Any charges for days beyond the long-stay outlier cutoff which are deemed not medically necessary shall be the responsibility of the beneficiary.

(g) * * *

(10) Amounts above allowable costs or charges. Costs of services and supplies to the extent amounts billed are over the CHAMPUS determined allowable cost or charge, as provided for in § 199.14.

(11) No legal obligation to pay, no charge would be made. Services or supplies for which the beneficiary or sponsor has no legal obligation to pay; or for which no charge would be made if the beneficiary or sponsor was not eligible under CHAMPUS, except claims subject to the CHAMPUS DRG-based payment system where the DRG-based amount is greater than the hospital's billed charge which has been paid in full by a double coverage plan.

4. Section 199.6 is amended by revising paragraphs (a)(8) and (b)(1)(ii), by adding a new paragraph (b)(3)(v), and by removing paragraph (e) and redesignating paragraph (f) as paragraph (e) to read as follows:

§ 199.6 Authorized providers.

* * *

(a) * * *

(8) Participating provider. Under CHAMPUS, authorized professional providers and institutional providers other than hospitals have the option of participating on a claim-by-claim basis. Participation is required for inpatient claims only for hospitals which are Medicare-participating providers. Hospitals which are not Medicare-participating providers but which are subject to the CHAMPUS DRG-based payment system in paragraph (a)(1)(ii)(D) of § 199.14 must sign agreements to participate on all CHAMPUS inpatient claims in order to

be authorized providers under CHAMPUS. All other hospitals may elect to participate on a claim-by-claim basis. Participating providers must indicate participation by signing the appropriate space on the applicable CHAMPUS claim form and submitting it to the appropriate CHAMPUS fiscal intermediary on behalf of the beneficiary. In the case of an institution or medical supplier, the claim must be signed by an official having such authority. This certifies that the provider has agreed to accept the CHAMPUSdetermined allowable charge or cost as payment in full for the medical services and supplies listed on the specific claim form; and has agreed to accept the amount paid by CHAMPUS or the CHAMPUS payment combined with the cost-sharing and deductible amounts paid by, or on behalf of, the beneficiary as full payment for the covered medical services and supplies. the terms to be a second

(b) * * * (1) * * *

(ii) Billing practices. Institutional billings must be itemized fully and sufficiently descriptive, even when CHAMPUS payment is determined under the CHAMPUS DRG-based payment system, so that CHAMPUS can make a determination of benefits. Except for claims subject to the CHAMPUS DRG-based payment system, whenever continuing charges are involved, claims should be submitted to the appropriate CHAMPUS fiscal intermediary at least every 30 days (monthly) either by the beneficiary or sponsor or directly by the provider on behalf of the beneficiary. For claims subject to the CHAMPUS DRG-based payment system, claims may be submitted only after the beneficiary has been discharged or transferred from the hospital.

(3) * * *

(v) Participation agreements required for some hospitals which are not Medicare-participating. Notwithstanding the provisions of this paragraph (B)(3), a hospital which is subject to the CHAMPUS DRG-based payment system but which is not a Medicare-particiating hospital must request and sign an agreement with OCHAMPUS. By signing the agreement, the hospital agrees to participate on all CHAMPUS inpatient claims and accept the requirements for a participating provider as contained in paragraph (a)(8) of § 199.6. Failure to sign such an agreement shall disqualify such hospital as a CHAMPUS-approved institutional provider.

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5. Section 199.7 is amended by revising paragraphs (B)(2)(i), (c)(2), (e)(1), and (g) and by adding a new paragraph (b)(2)(x)(C) to read as follows:

§ 199.7 Claims submissions, review, and payment.

(b) * * * (2) * * *

(i) Diagnosis. All applicable diagnoses are required; standard nomenclature is acceptable. In the absence of a diagnosis, a narrative description of the definitive set of symptoms for which the medical care was rendered must be provided.

(x) * * *

(C) For hospitals subject to the CHAMPUS DRG-based payment system (see paragraph (a)(1)(ii)(D) of § 199.14), the following information is also

(1) The principal diagnosis (the diagnosis established, after study, to be chiefly responsible for causing the patient's admission to the hospital).

(2) All secondary diagnoses.

(3) All significant procedures performed.

(4) The discharge status of the beneficiary.

(5) The hospital's Medicare provider number.

(6) The source of the admission. * * * * * * (c) * * *

(2) Provider's signature. A participating provider (see paragraph (a)(8) of § 199.6) is required to sign the CHAMPUS claim form.

(e) * * *

* *

(1) Continuing care. Except for claims subject to the CHAMPUS DRG-based payment system, whenever medical services and supplies are being rendered on a continuing basis, an appropriate claim or claims should be submitted every 30 days (monthly) whether submitted directly by the beneficiary or sponsor or by the provider on behalf of the beneficiary. Such claims may be submitted more frequently if the beneficiary or provider so elects. The Director, OCHAMPUS, or a designee, also may require more frequent claims submission based on dollars. Examples of care that may be rendered on a continuing basis are outpatient physical therapy, private duty (special) nursing, or inpatient stays. For claims subject to the CHAMPUS DRG-based payment

system, claims may be submitted only after the beneficiary has been discharged or transferred from the hospital.

(g) Claims review. It is the responsibility of the CHAMPUS fiscal intermediary (or OCHAMPUS, including OCHAMPUSEUR) to review each CHAMPUS claim submitted for benefit consideration to ensure compliance with all applicable definitions, conditions, limitations, or exclusions specified or enumerated in this part. It is also required that before any CHAMPUS benefits may be extended, claims for medical services and supplies will be subject to utilization review and quality assurance standards, norms, and criteria issued by the Director, OCHAMPUS, or a designee (see paragraph (a)(1)(v) of § 199.14 for review standards for claims subject to the CHAMPUS DRG-based payment system). * - *

6. Section 199.10 is amended by redesignating paragraph (a)(5)(iii) as paragraph (a)(5)(iv) and adding a new paragraph (a)(5)(iii) to read as follows:

§ 199.10 Appeal and hearing procedures.

(5) * * *

(iii) The establishment of diagnosisrelated groups (DRGs), or the methodology for the classification of inpatient discharges within the DRGs, or the weighting factors that reflect the relative hospital resources used with respect to discharges within each DRG. since each of these is established by this

7. A new § 199.14 is added to read as follows:

§ 199.14 Provider reimbursement methods.

(a) Hospitals. The CHAMPUSdetermined allowable cost for reimbursement of a hospital shall be determined on the basis of one of the following methodologies.

(1) CHAMPUS DRG-based payment system. Under the CHAMPUS DRGbased payment system, payment for the operating costs of inpatient hospital services furnished by hospitals subject to the system (generally short-term. acute-care hospitals) is made on the basis of prospectively determined rates and applied on a per discharge basis using Diagnosis Related Groups (DRGs). Payments under this system will include an urban/rural differentiation and an adjustment for area wage differences and indirect medical education costs. Additional payments will be made for

capital costs, direct medical education costs and outlier cases.

(i) General—(A) DRGs used. The CHAMPUS DRG-based payment system will use the same DRGs used in the most recently available grouper for the Medicare Prospective Payment System.

(B) Assignment of discharges to DRGs. (1) The classification of a particular discharge shall be based on the patient's age, sex, principal diagnosis (that is, the diagnosis established, after study, to be chiefly responsible for causing the patient's admission to the hospital), secondary diagnoses, procedures performed and discharge status.

(2) Each discharge shall be assigned to only one DRG regardless of the number of conditions treated or services furnished during the patient's stay.

(C) Basis of payment—(1) Hospital billing. Under the CHAMPUS DRGbased payment system, hospitals are required to submit claims (including itemized charges) in accordance with paragraph (b) of § 199.7. The CHAMPUS fiscal intermediary will assign the appropriate DRG to the claim based on the information contained on the claim.

(2) Payment on a per discharge basis. Under the CHAMPUS DRG-based payment system, hospitals are paid a predetermined amount per discharge for inpatient hospital services furnished to CHAMPUS beneficiaries.

(3) Claims priced as of date of discharge. All claims reimbursed under the CHAMPUS DRG-based payment system are to be priced as of the date of discharge, regardless of when the claim is submitted.

(4) Payment in full. The DRG-based amount paid for inpatient hospital services is the total CHAMPUS payment for the inpatient operating costs (as described in paragraph (a)(1)(i)(C)(5) of this section) incurred in furnishing services covered by the CHAMPUS. The full prospective payment amount is payable for each stay during which there is at least one covered day of care, except as provided in paragraph (a)(1)(iv)(D)(1)(i) of this section.

(5) Inpatient operating costs. The CHAMPUS DRG-based payment system provides a payment amount for inpatient operating costs, including:

(i) Operating costs for routine services; such as the costs of room, board, and routine nursing services;

(ii) Operating costs for ancillary services, such as hospital radiology and laboratory services (other than physicians' services) furnished to hospital inpatients;

(iii) Special care unit operating costs;

(iv) Malpractice insurance costs related to services furnished to inpatients.

(6) Discharges and transfers—(i) Discharges. A hospital inpatient is

discharged when:

(aa) The patient is formally released from the hospital (release of the patient to another hospital as described in paragraph (a)(1)(i)(C)(6)(I) (ii) of this section, or a leave of absence from the hospital, will not be recognized as a discharge for the purpose of determining payment under the CHAMPUS DRGbased payment system):
(bb) The patient dies in the hospital;

(cc) The patient is transerred from the care of a hospital included under the CHAMPUS DRG-based payment system to a hospital or unit that is excluded from the prospective payment system.

(ii) Transfers. Except as provided under paragraph (a)(1)(i)(C)(6)(i) of this section, a discharge of a hospital inpatient is not counted for purposes of the CHAMPUS DRG-based payment system when the patient is transferred:

(aa) From one patient area or unit of the hospital to another area or unit of

the same hospital:

(bb) From the care of a hospital included under the CHAMPUS DRGbased payment system to the care of another hospital paid under this system;

(cc) From the care of a hospital included under the CHAMPUS DRGbased payment system to the care of another hospital that is excluded from the CHAMPUS DRG-based payment system because of participation in a statewide cost control program which is exempt from the CHAMPUS DRG-based payment system under paragraph (a)(1)(ii)(A) of this section; or

(dd) From the care of a hospital included under the CHAMPUS DRGbased payment system to the care of a uniformed services treatment facility.

(iii) Payment in full to the discharging hospital. The hospital discharging an inpatient shall be paid in full under the CHAMPUS DRG-based payment system.

(iv) Payment to a hospital transferring an inpatient to another hospital. If a hospital subject to the CHAMPUS DRGbased payment system transfers an inpatient to another such hospital, the transferring hospital shall be paid a per diem rate, as determined under instructions issued by OCHAMPUS, for each day of the patient's stay in that hospital, not to exceed the DRG-based payment that would have been paid if the patient had been discharged to another setting. However, if a discharge is classified into DRG No. 385 (Neonates, died or transferred) or DRG

No. 456 (Burns, transferred to another acute care facility), the transferring hospital shall be paid in full.

(v) Additional payments to transferring hospitals. A transferring hospital may qualify for an additional payment for extraordinary cases that meet the criteria for long-stay or cost outliers.

(ii) Applicability of the DRG system-(A) Areas affected. The CHAMPUS DRG-based payment system shall apply to hospitals' services in the fifty states, the District of Columbia, and Puerto Rico, except that any State which has implemented a separate DRG-based payment system or similar payment system in order to control costs and is exempt from the Medicare Prospective Payment System may be exempt from the CHAMPUS DRG-based payment system if it requests exemption in writing, and provided payment under such system does not exceed payment which would otherwise be made under the CHAMPUS DRG-based payment system.

(B) Services subject to the DRG-based payment system. All normally covered inpatient hospital services furnished to CHAMPUS beneficiaries by hospitals are subject to the CHAMPUS DRG-

based payment system.

(C) Services exempt from the DRGbased payment system. The following hospital services, even when provided in a hospital subject to the CHAMPUS DRG-based payment system, are exempt from the CHAMPUS DRG-based payment system and shall be reimbursed under the procedures in paragraph (a)(2) of this section.

(1) Services provided by hospitals exempt from the DRG-based payment

system.

- (2) All services which would otherwise be paid under one of the psychiatric DRGs which are numbers 424-432.
- (3) All services which would otherwise be paid under one of the substance abuse DRGs which are numbers 433-438.
- (4) All services related to kidney acquisition by Renal Transplantation
- (5) All services related to a heart transplantation which would otherwise be paid under DRG 103.
- (6) All services related to liver transplantation when the transplant is performed in a CHAMPUS-authorized liver transplantation center.

(7) All professional services provided by hospital-based physicians.

(8) All services provided by nurse anesthetists.

(D) Hospitals subject to the CHAMPUS DRG-based payment system. All hospitals within the fifty States, the District of Columbia, and Puerto Rico which are certified to provide services to CHAMPUS beneficiaries are subject to the DRG-based payment system except for the following hospitals or hospital units which are exempt.

(1) Psychiatric hospitals. A
psychiatric hospital which is exempt
from the Medicare Prospective Payment
System is also exempt from the
CHAMPUS DRG-based payment
system. In order for a psychiatric
hospital which does not participate in
Medicare to be exempt from the
CHAMPUS DRG-based payment
system, it must meet the same criteria
(as determined by the Director,
OCHAMPUS, or a designee) as required
for exemption from the Medicare
Prospective Payment System as
contained in § 412.23 of Title 42 CFR.

(2) Rehabilitation hospitals. A rehabilitation hospital which is exempt from the Medicare Prospective Payment System is also exempt from the CHAMPUS DRG-based payment system. In order for a rehabilitation hospital which does not participate in Medicare to be exempt from the CHAMPUS DRG-based payment system, it must meet the same criteria (as determined by the Director, OCHAMPUS, or a designee) as required for exemption from the Medicare Prospective Payment System as contained in § 412.23 of Title 42 CFR.

(3) Alcohol/Drug hospitals. An alcohol/drug hospital which is exempt from the Medicare prospective payment system is also exempt from the CHAMPUS DRG-based payment system. In order for an alcohol/drug hospital which does not participate in Medicare to be exempt from the CHAMPUS DRG-based payment system, it must meet the same criteria (as determined by the Director, OCHAMPUS, or a designee) as required for exemption from the Medicare Prospective Payment System as contained in § 412.23 of Title 42 CFR.

(4) Psychiatric, rehabilitation and alcohol/drug units (distinct parts). A psychiatric, rehabilitation or alcohol/drug unit which is exempt from the Medicare prospective payment system is also exempt from the CHAMPUS DRG-based payment system. In order for a

distinct unit which does not participate in Medicare to be exempt from the CHAMPUS DRG-based payment system, it must meet the same criteria (as determined by the Director, OCHAMPUS, or a designee) as required for exemption from the Medicare Prospective Payment System as contained in § 412.23 of Title 42 CFR.

(5) Long-term hospitals. A long-term hospital which is exempt from the Medicare prospective payment system is also exempt from the CHAMPUS DRG-based payment system. In order for a long-term hospital which does not participate in Medicare to be exempt from the CHAMPUS DRG-based payment system, it must have an average length of inpatient stay greater than 25 days:

(i) As computed by dividing the number of total inpatient days (less leave or pass days) by the total number of discharges for the hospital's most recent fiscal year; or

(ii) As computed by the same method for the immediately preceding six-month period, if a change in the hospital's average length of stay is indicated.

(6) Sole community hospitals. Any hospital which has qualified for special treatment under the Medicare prospective payment system as a sole community hospital and has not given up that classification is exempt from the CHAMPUS DRG-based payment system. (See Subpart G of 42 CFR Part 412.)

(7) Christian Science sanitoriums. All Christian Science sanitoriums (as defined in paragraph (b)(4)(vii) of § 199.6) are exempt from the CHAMPUS DRG-based payment system.

(8) Cancer hospitals. Any hospital which qualifies as a cancer hospital under the Medicare standards and has elected to be exempt from the Medicare prospective payment system is exempt from the CHAMPUS DRG-based payment system. (See 42 CFR 412.94.)

(9) Children's hospitals. A children's hospital which is exempt from the Medicare Prospective Payment System is also exempt from the CHAMPUS DRG-based payment system. In order for a children's hospital which does not participate in Medicare to be exempt from the CHAMPUS DRG-based payment system, it must meet the same criteria (as determined by the Director, OCHAMPUS, or a designee) as required for exemption from the Medicare

Prospective Payment System as contained in 42 CFR 412.23.

(10) Hospitals outside the 50 states, the District of Columbia, and Puerto Rico. A hospital is excluded from the CHAMPUS DRG-based payment system if it is not located in one of the fifty States, the District of Columbia, or Puerto Rico.

(E) Hospitals which do not participate in Medicare. It is not required that a hospital be a Medicare-participating provider in order to be an authorized CHAMPUS provider. However, any hospital which is subject to the CHAMPUS DRG-based payment system and which otherwise meets CHAMPUS requirements but which is not a Medicare-participating provider (having completed a form HCFA-1514, Hospital Request for Certification in the Medicare/Medicaid Program and a form HCFA-1561, Health Insurance Benefit Agreement) must complete a participation agreement with OCHAMPUS. By completing the participation agreement, the hospital agrees to participate on all CHAMPUS inpatient claims and to accept the CHAMPUS-determined allowable amount as payment in full for these claims. Any hospital which does not participate in Medicare and does not complete a participation agreement with OCHAMPUS will not be authorized to provide services to CHAMPUS beneficiaries.

(iii) Determination of payment amounts. The actual payment for an individual claim under the CHAMPUS DRG-based payment system is calculated by multiplying the urban or rural adjusted standardized amounts (adjusted to account for area wage differences using the wage indexes used in the Medicare program) by a weighting factor specific to each DRG.

(A) Calculation of DRG Weights.

(1) Grouping of charges. All discharge records in the database shall be grouped by DRG.

(2) Remove DRGs 469 and 470. Records from DRGs 469 and 470 shall be removed from the database.

(3) Indirect medical education standardization. To standardize the charges for the cost effects of indirect medical education factors, each teaching hospital's charges will be divided by 1.0 plus the following ratio on a hospital-specific basis:

$$1.5 \times \left[\left| \text{E1.0+} \right| \begin{array}{c} \text{number of interns+residents} \\ \text{number of beds} \end{array} \right] .5795 - 1.0 \right]$$

(4) Wage level standardization. To standardize the charge records for area wage differences, each charge record will be divided into labor-related and nonlabor-related portions, and the labor-related portion shall be divided by the most recently available Medicare wage index for the area. The labor-related and nonlabor-related portions will then be added together.

(5) Elimination of statistical outliers.
All unusually high or low charges shall be removed from the database.

(6) Calculation of DRG average charge. After the standardization for indirect medical education, and area wage differences, an average charge for each DRG shall be computed by summing charges in a DRG and dividing that sum by the number of records in the DRG.

(7) Calculation of national average charge per discharge. A national average charge per discharge shall be calculated by summing all charges and dividing that sum by the total number of records from all DRG categories.

(8) DRG relative weights. DRG relative weights shall be calculated for each DRG category by dividing each DRG average charge by the national average charge.

(B) Empty and low-volume DRGs. The Medicare weight shall be used for any DRG with less than ten (10) occurrences in the CHAMPUS database. The short-stay thresholds shall be set at one day for these DRGs and the long-stay thresholds shall be set at the FY 87 Medicare thresholds.

(C) Updating DRG weights. The CHAMPUS DRG weights shall be updated or adjusted as follows:

(1) DRG weights shall be recalculated annually using CHAMPUS charge data and the methodology described in paragraph (a)(1)(iii)(A) of this section.

(2) When a new DRG is created, CHAMPUS will, if practical, calculate a weight for it using an appropriate charge sample (if available) and the methodology described in paragraph (a)(1)(iii)(A) of this section.

(3) In the case of any other change under Medicare to an existing DRG weight (such as in connection with technology changes), CHAMPUS shall adjust its weight for that DRG in a manner comparable to the change made by Medicare.

(D) Calculation of the adjusted standardized amounts. The following procedures shall be followed in calculating the CHAMPUS adjusted standardized amount.

(1) Differentiate urban and rural charges. All charges in the database shall be sorted into urban and rural groups. The following procedures will be applied to each group.

(2) Indirect medical education standardization. To standardize the charges for the cost effects of indirect medical education factors, each teaching hospital's charges will be divided by 1.0 plus the following ratio on a hospitalspecific basis:

(3) Wage level standardization. To standardize the charge records for area wage differences, each charge record will be divided into labor-related and nonlabor-related portions, and the labor-related portion shall be divided by the most recently available Medicare wage index for the area. The labor-related and nonlabor-related portions will then be added together.

(4) Apply the cost to charge ratio.

Each charge is to be reduced to a representative cost by using the Medicare cost to charge ratio increased by 1 percentage point in order to reimburse hospitals for bad debt expenses attributable to CHAMPUS beneficiaries. This results in an effective cost-to-charge ratio (adjusted for bad debt) of 0.67.

(5) Preliminary base year standardized amount. A preliminary base year standardized amount shall be calculated by summing all costs in the database applicable to the urban or rural group and dividing by the total

number of discharges in the urban or rural group.

(6) Update for inflation. The preliminary base year standardized amounts shall be updated using an annual update factor equal to 1.07 to produce fiscal year 1988 preliminary standardized amounts. Thereafter, development of a new standardized amount will use an inflation factor equal to the hospital market basket index used by the Health Care Financing Administration in their Prospective Payment System.

(7) The preliminary standardized amounts, updated for inflation, shall be divided by a system standardization factor so that total DRG outlays, given the database distribution across hospitals and diagnoses, are equal to the total charges reduced to costs.

(8) Labor and nonlabor portions of the adjusted standardized amounts. The adjusted standardized amounts shall be divided into labor and nonlabor portions in accordance with the Medicare division of labor and nonlabor portions.

(E) Adjustments to the DRG-based payment amounts. The following adjustments to the DRG-based amounts (the weight multiplied by the adjusted standardized amount) will be made.

(1) Outliers. CHAMPUS shall adjust the DRG-based payment to a hospital for atypical cases. These outliers are those cases that have either an unusually short length-of-stay or extremely long length-of-stay or that involve extraordinarily high costs when compared to most discharges classified in the same DRG.

(i) Length-of-stay outliers. Length-ofstay outliers shall be identified and paid by the fiscal intermediary when the claims are processed.

(aa) Short-stay outliers. Any discharge with a length-of-stay (LOS) less than 1.94 standard deviations from the DRG's geometric LOS shall be classified as a short-stay outlier. Short-stay outliers shall be reimbursed at 200 percent of the per diem rate for the DRG for each covered day of the hospital stay, not to exceed the DRG amount.

The per diem rate shall equal the DRG amount divided by the geometric mean length-of-stay for the DRG.

(bb) Long-stay outliers. Any discharge which has a length-of-stay (LOS) exceeding the lesser of 1.94 standard deviations or 17 days from the DRG's geometric LOS shall be classified as a long-stay outlier. Long-stay outliers shall be reimbursed the DRG-based amount plus 60 percent of the per diem rate for the DRG for each covered day of care beyond the long-stay outlier cutoff. The per diem rate shall equal the DRG amount divided by the geometric mean length-of-stay for the DRG.

(ii) Cost outliers. Any discharge which does not qualify as a length-of-stay outlier and which has standardized costs that exceed a threshold of the greater of two times the DRG-based amount or \$13,500 may be classified as a cost outlier. The standardized costs shall be calculated by multiplying the total charges by .67 and adjusting this amount for indirect medical education costs. Cost outliers shall be reimbursed the DRG-based amount plus 60 percent of all costs exceeding the threshold. Additional payment for cost outliers can be made only upon request by the hospital.

(2) Wage Adjustment. CHAMPUS will adjust the labor portion of the standardized amounts according to the

hospital's area wage index.

(3) Indirect Medical Education Adjustment. The wage adjusted DRG payment will also be multiplied by 1.0 plus the hospital's indirect medical education ratio.

(F) Updating the adjusted standardized amounts. Beginning in FY 1989, the adjusted standardized amounts will be updated by the Medicare annual update factor, unless the adjusted standardized amounts are recalculated.

(G) Annual Cost Pass-Throughs.

- (1) Capital costs. When requested in writing by a hospital, CHAMPUS shall reimburse the hospital its share of actual capital costs as reported annually to the CHAMPUS fiscal intermediary. Payment for capital costs shall be made annually based on the ratio of CHAMPUS inpatient days for those beneficiaries subject to the CHAMPUS DRG-based payment system to total inpatient days applied to the hospital's total allowable capital costs. Reductions in payments for capital costs which are required under Medicare shall also be applied to payments for capital costs under CHAMPUS.
- i) Costs included as capital costs. Allowable capital costs are those specified in Medicare Regulation § 413.130, as modified by § 412.72.

(ii) Services, facilities, or supplies provided by supplying organizations. If services, facilities, or supplies are provided to the hospital by a supplying organization related to the hospital within the meaning of Medicare Regulation § 413.17, then the hospital must include in its capital-related costs. the capital-related costs of the supplying organization. However, if the supplying organization is not related to the provider within the meaning of § 413.17. no part of the charge to the provider may be considered a capital-related cost unless the services, facilities, or supplies are capital-related in nature and:

(aa) The capital-related equipment is leased or rented by the provider;

(bb) The capital-related equipment is located on the provider's premises; and (cc) The capital-related portion of the

charge is separately specified in the

charge to the provider.

(2) Direct medical education costs. When requested in writing by a hospital, CHAMPUS shall reimburse the hospital its actual direct medical education costs as reported annually to the CHAMPUS fiscal intermediary. Such teaching costs must be for a teaching program approved under Medicare Regulation § 413.85. Payment for direct medical education costs shall be made annually based on the ratio of CHAMPUS inpatient days for those beneficiaries subject to the CHAMPUS DRG-based payment system to total inpatient days applied to the hospital's total allowable direct medical education costs. Allowable direct medical education costs are those specified in Medicare Regulation § 413.85.

(3) Information necessary for payment of capital and direct medical education costs. Any hospital subject to the CHAMPUS DRG-based payment system which wishes to be reimbursed for allowed capital and direct medical education costs must submit a report to the CHAMPUS fiscal intermediary. Such report is to be submitted within three months of the end of the hospital's Medicare cost-reporting period and shall cover the one-year period corresponding to the hospital's Medicare cost-reporting period. The first such report may cover a period of less than a full year—from the effective date of the CHAMPUS DRGbased payment system to the end of the hospital's Medicare cost-reporting period. All costs reported to the CHAMPUS fiscal intermediary must correspond to the costs reported on the hospital's Medicare cost report. (If these costs change as a result of a subsequent audit by Medicare, the revised costs are to be reported to CHAMPUS within 30 days of the date the hospital is notified of the change.) The report must be

signed by the hospital official responsible for verifying the amounts and shall contain the following information.

(i) The hospital's name. (ii) The hospital's address.

(iii) The hospital's CHAMPUS provider number.

(iv) The hospital's Medicare provider number.

(v) The period covered—this must correspond to the hospital's Medicare cost-reporting period.

(vi) Total inpatient days provided.

(vii) Total CHAMPUS inpatient days for those beneficiaries subject to the CHAMPUS DRG-based payment system provided.

(viii) Total allowable capital costs. (ix) Total allowable direct medical education costs.

(x) Total full-time equivalents for: (aa) Residents.

(bb) Interns.

(xi) Total inpatient beds as of the end of the cost-reporting period. If this has changed during the reporting period, an explanation of the change must be provided.

(xii) Title of official signing the report. (xiii) Reporting date.

(xiv) The report shall contain a certification statement that any changes to the items in paragraphs (a)(1)(iii)(G)(3) (vi), (vii), (viii), (ix), or (x), which are a result of an audit of the hospital's Medicare cost-report, shall be reported to CHAMPUS within thirty (30) days of the date the hospital is notified of the change.

(iv) Quality of care reviews.

(A) Objectives of review system. There are four required functions: (1) A review of the completeness,

adequacy and quality of care provided;

(2) A review of the reasonableness, necessity and appropriateness of hospital admissions under CHAMPUS DRG reimbursement;

(3) A validation of diagnoses and procedural information that determines CHAMPUS reimbursement; and

(4) A review of the necessity and appropriateness of care for which payment is sought on an outlier basis.

(B) Hospital cooperation. All hospitals which participate in CHAMPUS and submit CHAMPUS claims are required to provide all information necessary for CHAMPUS to properly process the claims. In order for CHAMPUS to be assured that services for which claims are submitted meet quality of care standards, hospitals are required to provide the peer review organization (PRO) responsible for quality review with all the information, within timeframes to be established by

OCHAMPUS, it needs to perform the review functions required by this paragraph. Additionally, all participating hospitals shall provide CHAMPUS beneficiaries, upon admission, with information about the admission and quality review system including their appeal rights. A hospital which does not cooperate in this activity shall be subject to termination as a CHAMPUS-authorized provider.

(C) Areas of review— (1) Admissions. The following areas shall be subject to review to determine whether inpatient care was medically appropriate and necessary, was delivered in the most appropriate setting and met acceptable standards of quality. This review may include preadmission or prepayment

review when appropriate.

(i) Transfers of CHAMPUS beneficiaries from a hospital or hospital unit subject to the CHAMPUS DRGbased payment system to another

hospital or hospital unit.

(ii) CHAMPUS admissions to a hospital or hospital unit subject to the CHAMPUS DRG-based payment system which occur within a certain period (specified by OCHAMPUS) of discharge from a hospital or hospital unit subject to the CHAMPUS DRG-based payment system.

(iii) A random sample of other CHAMPUS admissions for each hospital subject to the CHAMPUS DRG-based

payment system.

(iv) CHAMPUS admissions in any DRGs which have been specifically identified by OCHAMPUS for review or which are under review for any other reason.

- (2) DRG validation. The review organization responsible for quality of care reviews shall be responsible for ensuring that the diagnostic and procedural information reported by hospitals on CHAMPUS claims which is used by the fiscal intermediary to assign claims to DRGs is correct and matches the information contained in the medical records. In order to accomplish this, the following review activities shall be done.
- (i) Perform DRG validation reviews of each case under review.
- (ii) Review of claim adjustments submitted by hospitals which result in the assignment of a higher weighted DRG.
- (iii) Review for physician certification as to the major diagnoses and procedures and the physician's acknowledgment of annual receipt of the penalty statement as contained in the Medicare regulations at 42 CFR 412.40 and 412.46.
- (iv) Review of a sample of claims for each hospital reimbursed under the

CHAMPUS DRG-based payment system. Sample size shall be determined based upon the volume of claims submitted.

(3) Outlier review. Claims which qualify for additional payment as a long-stay outlier or as a cost-outlier shall be subject to review to ensure that the additional days or costs were medically necessary and appropriate and met all other requirements for CHAMPUS coverage. In addition, claims which qualify as short-stay outliers shall be reviewed to ensure that the admission was medically necessary and appropriate and that the discharge was not premature.

(4) Procedure review. Claims for procedures identified by OCHAMPUS as subject to a pattern of abuse shall be the subject of intensified quality

assurance review.

(5) Other review. Any other cases or types of cases identified by OCHAMPUS shall be subject to focused review.

(D) Actions as a result of review.—(1) Findings related to individual claims. If it is determined, based upon information obtained during reviews, that a hospital has misrepresented admission, discharge, or billing information, or is found to have quality of care defects, or has taken an action that results in the unnecessary admissions of an individual entitled to benefits, unnecessary multiple admission of an individual, or other inappropriate medical or other practices with respect to beneficiaries or billing for services furnished to beneficiaries, the entity responsible for admission and quality review in conjunction with the fiscal intermediary, shall, as appropriate:

(i) Recoup (in whole or in part) any amounts paid for the inpatient hospital services related to such an unnecessary admission or subsequent readmission and provide the hospital with a notice of

appeal rights; or

(ii) Require the hospital to take other corrective action necessary to prevent or correct the inappropriate practice.

(iii) Advise the hospital and beneficiary of appeal rights, as required by § 199.10 of this part.

(iv) Notify OCHAMPUS of all such actions.

- (2) Findings related to a pattern of inappropriate practices. In all cases where a pattern of inappropriate admissions and billing practices that have the effect of circumventing the CHAMPUS DRG-based payment system is identified, OCHAMPUS shall be notified of the hospital and practice involved.
- (3) Billed charges and set rates. The allowable costs for authorized care in

all hospitals not subject to the CHAMPUS DRG-based payment system shall be determined on the basis of billed charges or set rates. Under this procedure the allowable costs may not exceed the lower of:

(i) The actual charge for such service

made to the general public; or

(ii) The allowed charge applicable to the policyholders or subscribers of the CHAMPUS fiscal intermediary for comparable services under comparable circumstances, when extended to CHAMPUS beneficiaries by consent or agreement; or

(iii) The allowed charge applicable to the citizens of the community or state as established by local or state regulatory authority, excluding Title XIX of the Social Security Act or other welfare program, when extended to CHAMPUS beneficiaries by consent or agreement.

(b) Skilled Nursing Facilities (SNFs). The CHAMPUS-determined allowable cost for reimbursement of a SNF shall be determined on the same basis as for hospitals which are not subject to the CHAMPUS DRG-based payment

system.
(c) Reimbursement for Other Than Hospitals and SNFs. The Director, OCHAMPUS, or a designee, shall establish such other methods of determining allowable cost or charge reimbursement for those institutions, other than hospitals and SNFs, as may be required.

(d) Reimbursement of Freestanding Ambulatory Surgical Centers. Authorized care furnished by freestanding ambulatory surgical centers shall be reimbursed on the basis of the CHAMPUS-determined

reasonable cost.

- (e) Reimbursement of Individual Health-Care Professionals and Other Non-Institutional Health-Care Providers. The CHAMPUS-determined reasonable charge (the amount allowed by CHAMPUS) for the services of an individual health-care professional or other noninstitutional health-care provider (even if employed by or under contract to an institutional provider) shall be determined by one of the following methodologies, that is, whichever is in effect in the specific geographic location at the time covered services and supplies are provided to a CHAMPUS beneficiary.
- (1) Allowable charge method. The allowable charge method is the preferred and primary method for reimbursement of individual health-care professionals and other noninstitutional health-care providers.

(i) The allowable charge for authorized care shall be the lower of:

(A) The billed charge for the service;

(B) The prevailing charge level that does not exceed the amount equivalent to the 80th percentile of billed charges made for similar services in the same locality during the base period.

Note.—Public Law 97–86 provides that prevailing charges are to be determined at the 90th percentile. However, DOD Appropriation Acts have limited this to the 80th percentile. Prevailing charges shall continue to be calculated in accordance with any limitations set forth in the DOD Appropriation Acts, as implemented in instructions issued by the Director, OCHAMPUS.

(1) the 80th percentile of charges shall be determined on the basis of statistical data and methodology acceptable to the Director, OCHAMPUS, or a designee.

(2) The base period shall be a period of 12 calendar months and shall be adjusted at least once a year.

(ii) A charge that exceeds the prevailing charge can be determined to be allowable only when unusual circumstances or medical complications justify the higher charge. The allowable charge may not exceed the billed charge under any circumstances.

(2) Alternative method. The Director. OCHAMPUS, or a designee, may subject to the approval of the ASD(HA). establish an alternative method of reimbursement designed to produce reasonable control over health care costs and to ensure a high level of acceptance of the CHAMPUSdetermined charge by the individual health-care professionals or other noninstitutional health-care providers furnishing services and supplies to CHAMPUS beneficiaries. Alternative methods may not result in reimbursement greater than the allowable charge method above.

(f) Outside the United States. The Director, OCHAMPUS, or a designee, shall determine the appropriate reimbursement method or methods to be used in the extension of CHAMPUS benefits for otherwise covered medical services or supplies provided by hospitals or other institutional providers, physicians or other individual professional providers, or other providers outside the United States.

(g) Implementing Instructions. The Director, OCHAMPUS, or a designee. shall issue CHAMPUS policies, instructions, procedures, and guidelines, as may be necessary to implement the intent of this section.

Linda M. Lawson,

Alternate OSD Federal Register Liaison Officer, Department of Defense. August 24, 1987.

Table 1—CHAMPUS Weight and Threshold Summary

[Editorial Note: This table will not appear in the Code of Federal Regulations]

The following summary shows the final CHAMPUS DRG weights as well as arithmetic mean lengths of stay, geometric mean lengths of stay, and outlier thresholds. For those DRGs marked with an asterisk (i.e., the low volume DRGs), we substituted Medicare weights, length of stay values, and long stay thresholds. The short stay thresholds were set at one day for these DRGs.

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DRG		CHAMPUS	CHAMPUS ARITHMETIC GEOMETRIC SHORT STAY LONG STAY	GEOMETRIC	SHORT STAY	LONG STA	
NUMBER DESCRIPTION		WEIGHT	MEAN LOS	MEAN LOS	THRESHOLD THRESHOLD	THRESHOLI	
1 CRANIOTOMY AGE >=18 EXCEPT FOR TRAUMA	RAUMA	3.6060	13.3	10.2	2	72	
2 CRANIOTOMY FOR TRAUMA AGE >=18		4.4477	14.2	6.6	-	26	
3 CRANIOTOMY AGE <18		2.3836	9.5	6.2		23	
4 SPINAL PROCEDURES		2.1411	10.7	8.7	2	25	
5 EXTRACRANIAL VASCULAR PROCEDURES		1.7049	6.3	5.5	2	14	
6 CARPAL TUNNEL RELEASE		0.5402	2.3	1.8	-	9	
7 PERIPH & CRANIAL NERVE & OTHER NE	WERVE & OTHER NERV SYST PROC AGE >=70 &/OR C.C.	2.8290	11.4	7.1		72	
8 PERIPH & CRANIAL NERVE & OTHER NER	WERVE & OTHER WERV SYST PROC AGE <70 M/O C.C.	0.8608	3.8	2.8	-	12	
9 SPINAL DISORDERS & INJURIES		5.0299	11.4	4.9	-	23	
10 NERVOUS SYSTEM NEOPLASMS AGE >=70 AND/OR C.C.	0 AND/OR C.C.	1.2225	8.2	5.3	-	22	
11 NERVOUS SYSTEM NEOPLASMS AGE <70 W/O C.C	W/O C.C.	0.8829	5.9	3.8	-	20	511
12 DEGENERATIVE NERVOUS SYSTEM DISORDERS	RDERS	1.6168	12.1	6.7	-	23	
13 MULTIPLE SCLEROSIS & CEREBELLAR ATAXIA	ATAXIA	0.9771	7.6	5.0	-	21	1911/2
14 SPECIFIC CEREBROVASCULAR DISORDERS EXCEPT TIA	RS EXCEPT TIA	1.4611	8.2	5.7	-	22	
15 TRANSIENT ISCHEMIC ATTACKS AND PRECEREBRAL OCCLUSIONS	RECEREBRAL OCCLUSIONS	0.7454	4.1	3.2	うなか からい	12	
16 NONSPECIFIC CEREBROVASCULAR DISORDERS WITH C.C.	WERS WITH C.C.	1.2609	6.9	6.4	-	21	1
17 NONSPECIFIC CEREBROVASCULAR DISORDERS W/O C.C.	DERS W/O C.C.	0.7707	5.1	3.0	-	17	
18 CRANIAL & PERIPHERAL NERVE DISCRDERS AGE >=70 AND/OR C.C.	DERS AGE >=70 AND/OR C.C.	1.2039	7.9	5.5	-	22	
19 CRANIAL & PERIPHERAL NERVE DISORDERS AGE <70 W/O C.C.	DERS AGE <70 W/O C.C.	0.7152	5.4	3.8	100	19	
20 NERVOUS SYSTEM INFECTION EXCEPT VIRAL MENINGITIS	/IRAL MEMINGITIS	1.3990	8.5	0.9	-	22	NAME OF THE PARTY
21 VIRAL MENINGITIS		0.5790	4.1	3.5	-	11	
22 HYPERTENSIVE ENCEPHALOPATHY		0.7426	4.4	3.4	-	12	
23 NONTRAUMATIC STUPOR & COMA		0.9647	5.5	6.0	•	18	
24 SEIZURE & MEADACHE AGE >=70 AND/OR C.C.	% c.c.	0.9220	5.5	3.9	-	18	
	c.c.	0.5673	4.1	3.1	-	12	
26 SEIZURE & HEADACHE AGE 0-17		0.4713	3.2	2.4	-	6	
	KR	1.8402	7.2	3.9	-	20	100
	COMA, COMA <1 HR AGE >=70 AND/OR C.C.	1.2596	6.1	3.5	-	20	
29 TRAUMATIC STUPOR & COMA <1 HR AGE	COMA <1 HR AGE 18-69 W/O C.C.	0.6282	4.1	2.5	-	13	

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30 TRAUMATIC STUPOR & COMA <1 HR AGE 0-17 31 CONCUSSION AGE >=70 AND/OR C.C. 32 CONCUSSION AGE 18-69 W/O C.C. 33 CONCUSSION AGE 0-17 34 OTHER DISORDERS OF WERVOUS SYSTEM AGE >=70 AND/OR C.C. 35 OTHER DISORDERS OF NERVOUS SYSTEM AGE <70 W/O C.C. 36 RETINAL PROCEDURES 37 ORBITAL PROCEDURES 38 PRIMARY IRIS PROCEDURES 39 LENS PROCEDURES EXCEPT ORBIT AGE >=18 41 EXTRAOCULAR PROCEDURES EXCEPT ORBIT AGE 0-17 42 INTRAOCULAR PROCEDURES EXCEPT ORBIT AGE 0-17 43 HYPHEMA 44 ACUTE MAJOR EYE INFECTIONS		MEAN LOS H 2.6 4.1 2.4 7.9 4.2 3.1 3.1 1.9 1.7 3.0	1.8 1.9 1.9 1.3 5.0 2.8 2.9 2.9 2.5 1.7	THRESHOLD THRESHOLD 1	THRESHOLD 7 7 7 8 8 7 7 7 7 7 7 7 7 7 7 7 7 7 7
TRAUMATIC STUPOR CONCUSSION AGE 1 CONCUSSION AGE 0 OTHER DISORDERS OTHER DISORDERS OTHER DISORDERS OTHER DISORDERS OTHER DISORDERS EXTRACCULAR PROCI EXTRAOCULAR PROCI EXTRAOCULAR PROCI EXTRAOCULAR PROCI HYPHEMA ACUTE MAJOR EYE	0.4042 0.7241 0.3965 0.2338 1.7484 0.6492 0.8106 0.8111 0.4459 0.4851 0.4295	2.4.2.2.4.2.2.2.2.2.2.2.2.2.2.2.2.2.2.2	2.6 2.9 2.9 2.5 2.5 2.5 2.5 2.5 2.5		r 72 0 12 12 12 12 12 14 14 14 14 14 14 14 14 14 14 14 14 14
CONCUSSION AGE >> CONCUSSION AGE 11 CONCUSSION AGE 0 OTHER DISORDERS OTHER DISORDERS OTHER DISORDERS OTHER DISORDERS OTHER DISORDERS EXTRACCULAR PROCE EXTRACCULAR PROCE INTRACCULAR PROCE HYPHEMA ACUTE MAJOR EYE	0.7241 0.3965 0.2338 1.7484 0.6492 0.8111 0.4459 0.4851 0.4295	4.1 4.2 4.2 4.2 4.2 2.3 4.2 1.0 3.1 7.0 8.0 7.0 8.0 7.0 8.0 7.0 8.0 7.0 8.0 7.0 7.0 8.0 7.0 7.0 7.0 7.0 7.0 7.0 7.0 7.0 7.0 7	2.5 2.9 2.9 2.9 2.0 2.0 2.0 7.1		T 0 M 5 4 8 0 R 4 4
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CONCUSSION AGE O OTHER DISORDERS OTHER DISORDERS OTHER DISORDERS ORBITAL PROCEDUR ORBITAL PROCEDUR PRIMARY IRIS PROC LENS PROCULAR PROCI EXTRAOCULAR PROCI INTRAOCULAR PROCI HYPHEMA ACUTE MAJOR EYE	0.2338 1.7484 0.6492 0.8106 0.8111 0.4459 0.4851 0.4295	3.4 3.4 3.1 4.2 3.1 1.9 3.0 3.0 3.0	2.8 2.9 2.5 2.5 1.7 1.6		w 12 4 8 9 2 4 4
OTHER DISORDERS OTHER DISORDERS OTHER DISORDERS ORBITAL PROCEDURI ORBITAL PROCEDURI PRIMARY IRIS PROCI LENS PROCULAR PROCI EXTRAOCULAR PROCI INTRAOCULAR PROCI HYPHEMA ACUTE MAJOR EYE	1.7484 0.6492 0.8106 0.8111 0.4459 0.6959 0.4851 0.4295	2.2 2.2 2.2 2.2 2.2 7.1 3.0	5.0 2.9 2.5 2.0 1.7 1.6		2480844
RETINAL PROCEDURI ORBITAL PROCEDURI ORBITAL PROCEDURI PRIMARY IRIS PRO LENS PROCEDURES INTRAOCULAR PROCE INTRAOCULAR PROCE INTRAOCULAR PROCE HYPHEMA ACUTE MAJOR EYE	0.6492 0.8106 0.8111 0.4459 0.6959 0.4851 0.4295	4.2 3.4 2.2 2.2 1.9 3.0	2.9 2.5 2.5 2.0 1.7 1.6		480844
RETINAL PROCEDURES ORBITAL PROCEDURES PRIMARY IRIS PROCEDURES LENS PROCEDURES MITH OR MITHOL EXTRACCULAR PROCEDURES EXCEPT INTRACCULAR PROCEDURES EXCEPT INTRACCULAR PROCEDURES EXCEPT HYPHEMA ACUTE MAJOR EYE INFECTIONS	0.8106 0.8111 0.4459 0.6959 0.4851 0.4295	3.5 2.2 2.2 2.0 7.7 3.0	2.9 2.5 2.0 2.0 1.7		80 Or 10 4 A
ORBITAL PROCEDURES PRIMARY IRIS PROCEDURES LENS PROCEDURES WITH OR WITHOU EXTRAOCULAR PROCEDURES EXCEPT EXTRAOCULAR PROCEDURES EXCEPT INTRAOCULAR PROCEDURES EXCEPT HYPHEMA ACUTE MAJOR EYE INFECTIONS	0.8111 0.4459 0.6959 0.4851 0.4295	3.2 2.2 2.0 1.9 7.7	2.5 2.0 1.7		0 10 4 4
PRIMARY IRIS PROCEDURES LENS PROCEDURES WITH OR WITHOU EXTRAOCULAR PROCEDURES EXCEPT EXTRAOCULAR PROCEDURES EXCEPT INTRAOCULAR PROCEDURES EXCEPT HYPHEMA ACUTE MAJOR EYE INFECTIONS	0.4459 0.6959 0.4851 0.4295	2.2 2.0 1.9 1.7 3.0	1.7		N 4 4
LENS PROCEDURES WITH OR WITHOU EXTRAOCULAR PROCEDURES EXCEPT INTRAOCULAR PROCEDURES EXCEPT HYPHEMA ACUTE MAJOR EYE INFECTIONS	0.6959	1.9	1.7		4 4
EXTRAOCULAR PROCEDURES EXCEPT INTRAOCULAR PROCEDURES EXCEPT INTRAOCULAR PROCEDURES EXCEPT HYPHEMA ACUTE MAJOR EYE INFECTIONS	0.4851	1.9	1.6	-	7
EXTRAOCULAR PROCEDURES EXCEPT INTRAOCULAR PROCEDURES EXCEPT HYPHEMA ACUTE MAJOR EYE INFECTIONS	0.4295	1.7	7 6		-
INTRAOCULAR PROCEDURES EXCEPT HYPHEMA ACUTE MAJOR EYE INFECTIONS	0.7921	3.0	*-	-	3
43 HYPHEMA 44 ACUTE MAJOR EYE INFECTIONS			2.5	-	1
44 ACUTE MAJOR EYE INFECTIONS	0.3083	4.2	3.6	-	=
	0.4713	4.3	3.8	-	10
45 NEUROLOGICAL EYE DISORDERS	0.5503	3.1	2.6	-	80
46 OTHER DISORDERS OF THE EYE AGE >=18 WITH C.C.	0.6808	5.4	3.7	-	20
47 OTHER DISORDERS OF THE EYE AGE >=18 W/O C.C.	0.5982	4.4	3.0	-	15
48 OTHER DISORDERS OF THE EYE AGE 0-17	0.5002	3.8	5.6	-	12
49 MAJOR HEAD & NECK PROCEDURES	2.7560	11.6	8.6	-	25
50 SIALOADEMECTOMY	0.7656	5.6	2.1	-	9
51 SALIVARY GLAND PROCEDURES EXCEPT SIALOADENECTOMY	0.6647	5.9	2.3	1	0
52 CLEFT LIP & PALATE REPAIR	7769.0	3.4	5.9	-	80
53 SINUS & MASTOID PROCEDURES AGE >=18	0.7476	2.5	1.9	-	7
54 SINUS & MASTOID PROCEDURES AGE 0-17	0.6684	2.1	1.7	-	5
55 MISCELLANEOUS EAR, NOSE & THROAT PROCEDURES	0.4973	1.6	1.4	-	3
56 RHINOPLASTY	0.4822	1.6	1.4	-	3
	0.5202	2.5	2.0	-	9
58 T&A PROC EXCEPT TONSILLECTOMY &/OR ADENOIDECTOMY ONLY AGE 0-17	0.3662	1.3	1.2	-	2

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NUMBER DESCRIPTION	WE1GHT	CHAMPUS ARITHMETIC GEOMETRIC SHORT STAY LONG STAY WEIGHT MEAN LOS MEAN LOS THRESHOLD THRESHOLD	GEOMETRIC MEAN LOS	SHORT STAY LONG STAY THRESHOLD THRESHOLD	THRESHOLI
59 TONSILLECTOMY AND/OR ADENOIDECTOMY ONLY AGE >=18	0.3713	1.6	1.2	-	2
60 TONSILLECTOMY AND/OR ADENOIDECTOMY ONLY AGE 0-17	0.3357	1.3	1.2	-	2
	0.8733	2.8	1.9	-	10
62 MYRINGOTOMY WITH TUBE INSERTION AGE 0-17	0.3938	1.9	1.4		4
63 OTHER EAR, NOSE & THROAT O.R. PROCEDURES	0.9214	3.8	2.8	-	13
64 EAR, NOSE & THROAT MALIGNANCY	1.1117	6.2	3.9	-	20
65 DYSEQUILIBRIUM	0.5023	3.5	2.8	-	0
66 EPISTAXIS	0.5123	3.7	3.0	-	-
67 EPIGLOTTITIS	1,1043	4.3	3.6	-	12
68 OTITIS MEDIA & URI AGE >=70 AND/OR C.C.	0.6849	4.3	3.6	-	12
69 OTITIS MEDIA & URI AGE 18-69 W/O C.C.	0.4817	3.3	2.8	-	00
70 OTITIS MEDIA & URI AGE 0-17	0.4016	3.2	2.7	-	00
71 LARYNGOTRACHEITIS	0.3583	2.6	2.2		9
72 NASAL TRAUMA & DEFORMITY	0.3922	2.0	1.5		2
OTHER EAR, NOSE &	0.6875	3.7	2.6	-	12
	0.4760	3.0	2.1	-	0
	3.0582	11.4	9.6	M	92
	3.2224	12.0	7.9	-	72
	1.7591	7.3	5.0	-	22
78 PULMONARY EMBOLISM	1.6485	9.2	7.5	-	5%
79 RESPIRATORY INFECTIONS & INFLAMMATIONS AGE >=70 AND/OR C.C.	2.8086	11.0	8.6	-	10
80 RESPIRATORY INFECTIONS & INFLAMMATIONS AGE 18-69 W/O C.C.	1.6726	9.2	7.0	-	77
	1.3621	8.0	6.5	-	23
82 RESPIRATORY NEOPLASMS	1.3537	7.4	6.4	•	12
83 MAJOR CHEST TRAUMA AGE >=70 AND/OR C.C.	2.8879	4.6	4.9	-	23
MAJOR CHEST TRAUM	0.6126	4.4	3.5	-	13
	1.3375	7.4	5.5	-	22
	0.9337	6.1	4.1	-	12
87 PULMONARY EDEMA & RESPIRATORY FAILURE	2.6726	9.2	9.9	-	23

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DRG	CHAMPUS	ARITHMETIC	GEOMETRIC	CHAMPUS ARITHMETIC GEOMETRIC SHORT STAY LONG STAY	IG STAY
NUMBER DESCRIPTION	WEIGHT	MEAN LOS	MEAN LOS	MEAN LOS THRESHOLD THRESHOLD	RESHOLD
88 CHRONIC OBSTRUCTIVE PULMONARY DISEASE	1.3593	6.9	5.4	-	12
89 SIMPLE PNEUMONIA & PLEURISY AGE >=70 AND/OR C.C.	1.5290	7.5	6.3	-	20
90 SIMPLE PNEUMONIA & PLEURISY AGE 18-69 W/O C.C.	0.9350	5.5	4.7		14
91 SIMPLE PHEUMONIA & PLEURISY AGE 0-17	0.6261	4.3	3.6	-	10
92 INTERSTITIAL LUNG DISEASE AGE >=70 AND/OR C.C.	1.5426	8.5	5.3	-	22
93 INTERSTITIAL LUNG DISEASE AGE <70 W/O C.C.	0.8010	4.7	3.7	-	15
	1.5808	8.6	6.1		23
95 PNEUMOTHORAX AGE <70 W/O C.C.	0.6552	5.0	4.1	-	14
96 BRONCHITIS & ASTHMA AGE >=70 AND/OR C.C.	1.1797	6.2	5.2	1	16
97 BRONCHITIS & ASTHMA AGE 18-69 W/O C.C.	0.8014	4.8	4.1		12
98 BRONCHITIS & ASTHMA AGE 0-17	0.5031	3.4	2.9		80
99 RESPIRATORY SIGNS & SYMPTOMS AGE >=70 AND/OR C.C.	0.9411	8.4	3.7		15
100 RESPIRATORY SIGNS & SYMPTOMS AGE <70 W/O C.C.	0.5788	3.3	2.5	-	6
OTHER RESPIRATORY	1.3947	6.2	4.5	-	12
	0.7468	3.9	2.8	-	13
	9762.9	12.9	11.3	n	28
	5.8168	10.7	6.5	2	92
	5.8104	12.0	11.0	7	77
	4.8904	10.0	9.5	7	12
	3.7739	10.3	8.6	2	22
OTHER CARDIOTHORA	4.0747	10.7	8.6	2	25
MAJOR RECONSTRUCT	3.7060	11.7	6.6	3	56
MAJOR RECONSTRUCT	2.7987	9.3	8.4	2	20
	2,3117	8.9	5.1		12
113 AMPUTATION FOR CIRC SYSTEM DISORDERS EXCEPT UPPER LIMB & TOE	3,6268	19.8	15.1	n	32
114 UPPER LIMB & TOE AMPUTATION FOR CIRC SYSTEM DISORDERS	2.0505	11.5	8.8	-	25
	4.7176	11.4	9.1	-	92
116 PERMANENT CARDIAC PACEMAKER IMPLANT W/O AMI, HEART FAILURE OR SHO	3.1991	0.9	5.0	•	16

CO	CHAMPUS	CHAMPUS ARITHMETIC GEOMETRIC SHORT STAY LONG STAY	GEOMETRIC	SHORT STAY	LONG STAY
NUMBER DESCRIPTION	WEIGHT	MEAN LOS	MEAN LOS	MEAN LOS THRESHOLD THRESHOLD	THRESHOLD
117 CARDIAC DACEMAKER REDIACE & REVIS EXC PULSE GEN REPL	1.4970	6.2	4.1		12
118 CARDIAC PACEMAKER PILSE GENERATOR REPLACEMENT	2.1962	2.7	2.3	-	7
110 VEIM LIGATION & STRIPPING	0.7846	3.7	3.0	-	10
120 OTHER O.R. PROCEDURES ON THE CIRCULATORY SYSTEM	2.8012	12.5	7.1	-	57
121 CIRCULATORY DISORDERS WITH AMI & C.V. COMP. DISCH. ALIVE	2.1709	8.8	6.9	-	23
122 CIRCULATORY DISORDERS WITH AMI W/O C.V. COMP. DISCN. ALIVE	1,5068	6.5	5.2	-	22
123 CIRCULATORY DISORDERS WITH AMI, EXPIRED	2.3655	4.2	2.6	-	16
124 CIRCULATORY DISORDERS EXC AMI, WITH CARD CATH & COMPLEX DIAG	1.4518	8.4	3.7	-	15
125 CIRCULATORY DISORDERS EXC AMI, WITH CARD CATH W/O COMPLEX DIAG	7698.0	3.0	2.3	-	8
126 ACUTE & SUBACUTE ENDOCARDITIS	3.7074	17.8	12.7	2	53
	1.2739	6.5	5.1	-	20
128 DEEP VEIN THROMBOPHLEBITIS	0.9339	7.8	6.7	2	20
129 CARDIAC ARREST, UNEXPLAINED	3.5298	10.4	4.7	-	21
-	1.1482	7.0	6.4	-	21
131 PERIPHERAL VASCULAR DISORDERS AGE <70 W/O C.C.	0.7643	5.2	3.7	-	19
132 ATHEROSCLEROSIS AGE >=70 AND/OR C.C.	1.0702	3.6	2.8	-	10
ATHEROSCLEROSIS	0.9704	3.2	2.6		6
	0.6540	4.3	3.4	-	12
135 CARDIAC CONGENITAL & VALVULAR DISORDERS AGE >=70 AND/OR C.C.	0.9561	3.9	3.1		11
136 CARDIAC CONGENITAL & VALVULAR DISORDERS AGE 18-69 W/O C.C.	0.7412	3.2	2.5		6
137 CARDIAC CONGENITAL & VALVULAR DISORDERS AGE 0-17	0.7581		2.1		0
138 CARDIAC ARRHYTHMIA & CONDUCTION DISORDERS AGE >=70 AND/OR C.C.	0.8866	4.4	3.4		13
139 CARDIAC ARRHYTHMIA & CONDUCTION DISORDERS AGE <70 W/O C.C.	0.6432		2.7		10
140 ANGINA PECTORIS	0.8099	3.6	3.0		10
141 SYNCOPE & COLLAPSE AGE >=70 AND/OR C.C.	0.6900	0.4	3.1		12
142 SYNCOPE & COLLAPSE AGE <70 W/O C.C.	0.5358	3.0	2.4		8
143 CHEST PAIN	96290	2.9	2.4		7
144 OTHER CIRCULATORY DIAGNOSES MITH C.C.	1.3249	6.3	4.5		12
145 OTHER CIRCULATORY DIAGNOSES M/O C.C.	0.8308	4.1	3.1		13

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DRG MINIERE DECEMBATION	CHAMPUS	ARITHMETIC	GEOMETRIC	CHAMPUS ARITHMETIC GEOMETRIC SHORT STAY LONG STAY	LONG STA
NOTICE VESCRIPTION	WEIGHT	MEAN LOS	MEAN LOS	MEAN LOS THRESHOLD THRESHOLD	THRESHOL
146 RECTAL RESECTION AGE >=70 AND/OR C.C.	3.2885	14.4	13.0	5	&
147 RECTAL RESECTION AGE <70 W/O C.C.	2.3506	10.8	10.3	5	18
148 MAJOR SMALL & LARGE BOWEL PROCEDURES AGE >=70 AND/OR C.C.	3.5614	13.9	11.9	7	28
149 MAJOR SMALL & LARGE BOWEL PROCEDURES AGE <70 W/O C.C.	2.0799	10.1	8.8	~	24
150 PERITOWEAL ADMESTOLYSIS AGE >=70 AND/OR C.C.	2.3009	11.1	9.8	2	25
151 PERITONEAL ADHESIOLYSIS AGE <70 W/O C.C.	1.5484	8.2	7.2	2	20
152 MINOR SMALL & LARGE BOWEL PROCEDURES AGE >=70 AND/OR C.C.	2.2572	10.01	7.9	2	72
153 MINOR SMALL & LARGE BOWEL PROCEDURES AGE <70 W/O C.C.	1.3399	7.5	6.2	-	21
154 STOMACH, ESOPHAGEAL & DUODENAL PROCEDURES AGE >=70 AND/OR C.C.	3.4536	12.3	10.0	2	26
155 STOMACH, ESOPHAGEAL & DUODENAL PROCEDURES AGE 18-69 W/O C.C.	2.0912	9.1	7.6	2	24
156 STOMACH, ESOPHAGEAL & DUCDENAL PROCEDURES AGE 0-17	1.0770	5.6	4.3	-	16
157 AMAL AND STOMAL PROCEDURES AGE >=70 AND/OR C.C.	0.9817	5.6	4.5	-	17
158 ANAL AND STOMAL PROCEDURES AGE <70 W/O C.C.	0.6166	3.7	3.1	-	10
159 HERNIA PROCEDURES EXCEPT INGUINAL & FEMORAL AGE >=70 AND/OR C.C.	1.2607	0.9	5.0	-	16
160 HERNIA PROCEDURES EXCEPT INGUINAL & FEMORAL AGE 18-69 W/O C.C.	0.8122	0.4	3.3	-	=
161 INGUINAL & FEMORAL HERNIA PROCEDURES AGE >=70 AND/OR C.C.	0.8475	3.9	3.2	-	1
	0.5421	2.4	2.0	-	9
163 HERNIA PROCEDURES AGE 0-17	0.4304	1.8	1.5	-	7
164 APPENDECTOMY WITH COMPLICATED PRINC. DIAG AGE>=70 AND/OR C.C.	2.3864	10.1	9.3	7	20
165 APPENDECTOMY WITH COMPLICATED PRINC. DIAG AGE <70 W/O C.C.	1.2534	6.3	5.6	2	15
166 APPENDECTOMY W/O COMPLICATED PRINC. DIAG AGE >=70 AND/OR C.C.	1.2428	5.9	5.1	-	16
167 APPENDECTOMY W/O COMPLICATED PRINC. DIAG AGE <70 W/O C.C.	8669.0	3.6	3.2	-	7
168 PROCEDURES ON THE MOUTH AGE >=70 AND/OR C.C.	1.8458	4.9	9.4	-	21
169 PROCEDURES ON THE MOUTH AGE <70 U/O C.C.	0.9303	3.2	2.7	-	80
170 OTHER DIGESTIVE SYSTEM O.R. PROCEDURES AGE >=70 AND/OR C.C.	2.9384	13.6	6.6	-	26
171 OTHER DIGESTIVE SYSTEM O.R. PROCEDURES AGE <70 W/O C.C.	1.2247	0.9	4.4	-	21
172 DIGESTIVE MALIGNANCY AGE >=70 AND/OR C.C.	1.4151	8.6	5.6	-	22
173 DIGESTIVE MALIGNANCY AGE <70 W/O C.C.	1.0248	6.2	4.1	-	21
174 G.I. MEMORRHAGE AGE >=70 AND/OR C.C.	1.1540	5.7	4.4	-	17

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DRG	CHAMPUS	CHAMPUS ARITHMETIC GEOMETRIC SHORT STAY LONG STAY	GEOMETRIC	SHORT STAY	LONG STAY
NUMBER DESCRIPTION	WEIGHT	MEAN LOS	MEAN LOS	MEAN LOS THRESHOLD THRESHOLD	THRESHOLD
175 G.1. HEMORRHAGE AGE <70 W/O C.C.	0.7029	4.2	3.5	-	11
176 COMPLICATED PEPTIC ULCER	0.9471	5.3	4.2	-	16
177 UNCOMPLICATED PEPTIC ULCER >=70 AND/OR C.C.	0.8815	5.4	9.4	-	13
178 UNCOMPLICATED PEPTIC ULCER <70 W/O C.C.	0.6263	4.2	3.5	-	=
179 INFLAMMATORY BOWEL DISEASE	1.0332	7.0	5.3	-	22
180 G.I. OBSTRUCTION AGE >=70 AND/OR C.C.	1.0272	9.9	6.4	-	12
181 G.1. OBSTRUCTION AGE <70 W/O C.C.	0.5976	4.3	3.5	-	12
182 ESOPHAGITIS, GASTROENT.& MISC. DIGEST. DIS AGE >=70 %/OR C.C.	0.7497	4.8	3.8	-	11
183 ESOPHAGITIS, GASTROENT. & MISC. DIGEST. DIS AGE 18-69 W/O C.C.	0.5574	3.8	3.0	-	-
184 ESOPHAGITIS, GASTROENTERITIS & MISC. DIGEST. DISORDERS AGE 0-17	0.3297	2.9	2.4	-	1
185 DENTAL & ORAL DIS. EXC EXTRACTIONS & RESTORATIONS, AGE >=18	0.8997	4.8	3.3	-	17
186 DENTAL & ORAL DIS. EXC EXTRACTIONS & RESTORATIONS, AGE 0-17	0.4184	3.0	2.5		80
187 DENTAL EXTRACTIONS & RESTORATIONS	0.4942	2.1	1.7	-	
188 OTHER DIGESTIVE SYSTEM DIAGNOSES AGE >=70 AND/OR C.C.	1.0015	5.4	4.1	-	17
189 OTHER DIGESTIVE SYSTEM DIAGNOSES AGE 18-69 M/O C.C.	0.6098	0.4	2.8	-	13
190 OTHER DIGESTIVE SYSTEM DIAGNOSES AGE 0-17	0.4015	2.6	1.9	-	7
191 MAJOR PANCREAS, LIVER & SHUNT PROCEDURES	5.0848	16.4	13.3	M	30
192 MINOR PANCREAS, LIVER & SHUNT PROCEDURES	5.2655	18.7	12.6	2	29
193 BILIARY TRACT PROC EXC TOT CHOLECYSTECTOMY AGE >=70 &/OR C.C.	3.3429	13.7	10.7	2	27
194 BILIARY TRACT PROC EXC TOT CHOLECYSTECTOMY AGE <70 W/O C.C.	2.0402	9.6	4.8	2	54
195 TOTAL CHOLECYSTECTOMY WITH C.D.E. AGE >= 70 AND/OR C.C.	2.0491	4.6	8.7	*	18
196 TOTAL CHOLECYSTECTOMY WITH C.D.E. AGE <70 W/O C.C.	1.4977	7.9	7.3	2	15
197 TOTAL CHOLECYSTECTOMY W/O C.D.E. AGE >=70 AND/OR C.C.	1.4689	7.4	6.7	2	15
198 TOTAL CHOLECYSTECTOMY W/O C.D.E. AGE <70 W/O C.C.	1.0987	5.8	5.4	2	=
199 HEPATOBILIARY DIAGNOSTIC PROCEDURE FOR MALIGNANCY	1.9111	10.9	7.6	M	26
200 MEPATOBILIARY DIAGNOSTIC PROCEDURE FOR MON-MALIGNANCY	2.4732	10.0	8.9	-	23
	2.7311	10.2	7.0	-	77
202 CIRRHOSIS & ALCOHOLIC HEPATITIS	1.3853	7.7	5.5	-	22
203 MALIGNANCY OF NEPATOBILIARY SYSTEM OR PANCREAS	1,2989	7.9	5.3	-	22

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204 DISORDERS OF PANCREAS EXCEPT MALIGNANCY	1.0126	6.5	5.2	1	18
205 DISORDERS OF LIVER EXC MALIG, CIRR, ALC HEPA AGE >=70 AND/OR C.C.	1.7588	8.7		-	23
DISORDERS OF LIVE	0.7416	5.0			18
207 DISORDERS OF THE BILIARY TRACT AGE >=70 AND/OR C.C.	0.9392	5.3		-	17
208 DISORDERS OF THE BILIARY TRACT AGE <70 M/O C.C.	0.6001	3.5		-	10
	2.9943	11.4		5	22
	3.2031	14.6		3	28
	1.9295	9.8		2	25
212 HIP & FEMUR PROCEDURES EXCEPT MAJOR JOINT AGE 0-17	1.4129	8.7		-	22
	1.7263	10.7		-	77
	2.4274	11.1	9.3	2	26
215 BACK & NECK PROCEDURES AGE <70 W/O C.C.	1.5126	7.9	6.8	2	19
216 BIOPSIES OF MUSCULOSKELETAL SYSTEM & CONNECTIVE TISSUE	1.3089	6.6		-	21
	2.0273	6.6	5.5	-	22
218 LOWER EXTREM & HUNER PROC EXC HIP, FOOT, FENUR AGE >=70 &/OR C.C.	1.8658	8.8	7.1	-	24
219 LOWER EXTREM & HUMER PROC EXC HIP, FOOT, FEMUR AGE 18:69 W/O C.C.	1.0855	5.1	4.2	-	16
220 LOWER EXTREM & HUMER PROC EXC NIP, FOOT, FEMUR AGE 0-17	0.8099	3.6	2.6	-	11
221 KNEE PROCEDURES AGE >=70 AND/OR C.C.	1.9419	6.6	4.1	-	21
222 KNEE PROCEDURES AGE <70 M/O C.C.	0.8425	3.4	2.7	-	6
MAJOR SHOULDER/ELE	1.1151	4.5	3.4	-	14
	0.7700	2.9	2.4	-	7
	0.7177	3.0	2.5	-	80
	1.0876	4.9	4.3	-	21
SOFT TISSUE PROCED	0.7236	3.4	2.5	-	6
	0.8723	3.2	2.6	•	6
	0.6455	2.3	1.8	-	9
LOCAL EXCISION & REMOVAL OF INT	0.7155	3.6	2.3	-	11
231 LOCAL EXCISION & REMOVAL OF INT FIX DEVICES EXCEPT HIP & FEMUR	0.8753	3.8	2.6	-	12
232 ARTHROSCOPY	0,6393	2.3	1.8	-	9

DRG		CHAMPUS	ARITHMETIC	CHAMPUS ARITHMETIC GEOMETRIC SHORT STAY LONG STAY	SHORT STAY	LONG STAY	
NUMBER	NUMBER DESCRIPTION	WEIGHT	MEAN LOS	MEAN LOS	MEAN LOS THRESHOLD THRESHOLD	THRESHOLD	
233	OTHER MUSCULOSKELET SYS & CONN TISS O.R. PROC AGE >=70 &/OR C.C.	2.4202	9.7	7.2	-	54	
234	OTHER MUSCULOSKELET SYS & CONN	0.9855	4.7			13	
235	FRACTURES OF FEMUR	1.2545	13.7		-	77	
236	FRACTURES OF HIP & PELVIS	0.9995	8.8		-	23	
237	SPRAINS, STRAINS, & DISLOCATIONS OF MIP, PELVIS & THIGH	0.6421	5.0			16	
238	OSTEOMYELITIS	1.6239	11.9		-	25	
239	PATHOLOGICAL FRACTURES & MUSCULOSKELETAL & CONN.TISS.MALIGNANCY	1.1907	7.8		-	23	
240		1.4885	8.4		•	23	
241	CONNECTIVE TISSUE DISORDERS AGE <70 W/O C.C.	0.8490	6.5		-	21	
242	SEPTIC ARTHRITIS	1.2112	9.6		2	54	
243	MEDICAL BACK PROBLEMS	0.7425	5.6		-	20	
544	BONE DISEASES & SPECIFIC ARTHROPATHY AGE >=70 AND/OR C.C.	0.8232	6.0		•	21	V S
245	BONE DISEASES & SPECIFIC ARTHROPATHY AGE <70 W/O C.C.	0.6544	4.8		•	17	
546	246 NON-SPECIFIC ARTHROPATHIES	0.5751	3.8	3.2	•	10	
247	SIGNS & SYMPTOMS OF MUSCULOSKELETAL SYSTEM & CONN TISSUE	0.6729	5.2	3.5	-	19	
248	TENDONITIS, MYOSITIS & BURSITIS	0.6042	4.3		-	13	
549	249 AFTERCARE, MUSCULOSKELETAL SYSTEM & CONNECTIVE TISSUE	0.7924	5.4	3.1	-	20	
250	FX, SPRNS, STRNS & DISL OF FOREARM, HAND, FOOT AGE >=70 &/OR C.C.	0.7355	5.1		-	17	
251	251 FX, SPRNS, STRNS & DISL OF FOREARM, HAND, FOOT AGE 18-69 W/O C.C.	0.4550	2.7		-	7	THE STREET
252	FX,SPRNS,STRNS & DISL OF FOREARM, MAND, FOOT AGE 0-17	0.3186	1.4		-	2	
253	FX, SPRNS, STRNS & DISL OF UPARM, LOWLEG EX FOOT AGE >=70 &/OR C.C.	0.8188	5.5		-	18	
524	FX, SPRNS, STRNS & DISL OF UPARM, LOWLEG EX FOOT AGE 18-69 W/O C.C.	0.4732	3.4	2.6	-	10	
255	FX, SPRNS, STRNS & DISL OF UPARM, LOWLEG EX FOOT AGE 0-17	0.3709	2.5	1.8	-	9	
256	OTHER DIAGNOSES OF MUSCULOSKELETAL SYSTEM & CONNECTIVE TISSUE	9099.0	4.1		•	13	
257	TOTAL MASTECTOMY FOR MALIGNANCY AGE >=70 AND/OR C.C.	1.2098	6.1	5.5	2	13	
258	TOTAL MASTECTOMY FOR MALIGNANCY AGE <70 W/O C.C.	1.0060	5.0		-	11	
259	SUBTOTAL MASTECTOMY FOR MALIGNANCY AGE >=70 AND/OR C.C.	1.1880	7.0		-	12	
260	260 SUBTOTAL MASTECTOMY FOR MALIGNANCY AGE <70 W/O C.C	0.7876	3.2	2.4	-	0	
261	261 BREAST PROC FOR NON-MALIG EXCEPT BIOPSY & LOC EXC	0.9072	3.2	77	· Control of	×	1 =

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DRG	CHAMPUS	CHAMPUS ARITHMETIC GEOMETRIC SHORT STAY LONG STAY	GEOMETRIC	SHORT STAY	LONG STAY
NUMBER DESCRIPTION	WEIGHT	MEAN LOS	MEAN LOS	THRESHOLD	THRESHOLD THRESHOLD
262 BREAST BIOPSY & LOCAL EXCISION FOR NON-MALIGNANCY	0.5166	2.1	8.1		5
263 SKIN GRAFTS &/OR DEBRID FOR SKIN ULCER OR CELLULITIS AGE>=70 &/O	2.9552	16.5	12.0	2	28
264 SKIN GRAFTS &/OR DEBRID FOR SKIN ULCER OR CELLULITIS AGE <70 W/O	1.8731	12.0	9.0	•	25
265 SKIN GRAFTS &/OR DEBRID EXCEPT FOR SKIN ULCER OR CELLULITIS WITH	1.7366	7.6	6.9	-	23
266 SKIN GRAFTS &/OR DEBRID EXCEPT FOR SKIN ULCER OR CELLULITIS W/O	0.9241	4.7	3.3	-	16
267 PERIANAL & PILONIDAL PROCEDURES	0.4913	2.4	2.0	-	9
268 SKIN, SUBCUTANEOUS TISSUE & BREAST PLASTIC PROCEDURES	0.7085	2.9	2.4	-	80
269 OTHER SKIN, SUBCUT TISS & BREAST O.R. PROC AGE >=70 &/OR C.C.	1.5437	7.7	5.5	-	22
	0.7451	3.6	2.5	-	12
271 SKIN ULCERS	1.1258	8.3	6.5	-	23
272 MAJOR SKIN DISORDERS AGE >=70 AND/OR C.C.	1.1249	8.3	5.7	-	22
273 MAJOR SKIN DISORDERS AGE <70 W/O C.C.	0.7077	6.1	4.4	-	12
274 MALIGNANT BREAST DISORDERS AGE >=70 AND/OR C.C.	1.2667	8.8	6.0	-	22
	0.7625	5.4	3.7	-	20
	0.5332	3.1	2.6	-	80
277 CELLULITIS AGE >=70 AND/OR C.C.	1.0255	8.9	5.7	-	17
	0.7518	5.3	4.5	-	14
279 CELLULITIS AGE 0-17	0.5277	4.3	3.5	-	11
	0.5775	3.7	2.7	-	12
	0.4130	2.8	2.2	-	80
	0.3181	2.2	1.7	-	2
	0.7153	6.4	4.1	•	13
284 MINOR SKIN DISORDERS AGE <70 W/O C.C.	0.5054	3.8	5.9	-	11
285 AMPUTATION OF LOWER LIMB FOR ENDOCRINE, NUTRITIONAL & METABOLIC D	2.3351	15.4	14.0	5	30
	2.8083	10.6	8.9	2	52
287 SKIN GRAFIS & WOUND DEBRIDE FOR ENDOC, NUTRIT & METAB DISORDERS	1.9411	12.4	10.0	2	27
288 O.R. PROCEDURES FOR OBESITY	1.8201	6.5	6.1	2	12
289 PARATHYROID PROCEDURES	0.9714	9.4	4.0	-	10
290 THYROID PROCEDURES	0.7745	3.2	2.8	-	1

CHAMPUS WEIGHT AND THRESHOLD SUMMARY

DRG		CHAMPUS	ARITHMETIC	GEOMETRIC	CHAMPUS ARITHMETIC GEOMETRIC SHORF STAY LONG STAY	LONG STAY
NUMBER DESCRIPTION	CRIPTION	WEIGHT	MEAN LOS	MEAN LOS	MEAN LOS THRESHOLD THRESHOLD	THRESHOLD
291 THY	THYROGLOSSAL PROCEDURES	0.4241	1.6	1.6		23
292 OTHI	OTHER ENDOCRINE, NUTRIT & METAB O.R. PROC AGE >70 &/OR C.C.	2.0491	9.0	7.6	2	54
293 OTHI	OTHER ENDOCRINE, NUTRIT & METAB O.R. PROC AGE <70 W/O C.C.	1.5269	4.9	6.4	-	21
294 DIA	DIABETES AGE >=36	0.7556	5.0	5.0	-	15
295 DIA	DIABETES AGE 0-35	0.6385	4.7	3.9	-	13
	NUTRITIONAL & MISC. METABOLIC DISORDERS AGE >=70 AND/OR C.C.	1.0660	4.9	4.8	-	21
	NUTRITIONAL & MISC. METABOLIC DISORDERS AGE 18-69 W/O C.C.	0.7334	5.2	3.7	-	17
	NUTRITIONAL & MISC. METABOLIC DISORDERS AGE 0-17	0.6039	8.4	3.4	-	16
	INBORN ERRORS OF METABOLISM	1.0303	5.6	3.9	-	19
	ENDOCRINE DISORDERS AGE >=70 AND/OR C.C.	1.0597	6.2	4.8	-	20
301 END	ENDOCRINE DISORDERS AGE <70 W/O C.C.	0.6648	9.4	3.1	-	15
	KIDNEY TRANSPLANT	3.7135	17.5	15.2	5	32
303 KID	KIDNEY, URETER & MAJOR BLADDER PROCEDURE FOR NEOPLASM	2.3414	10.3	9.5	7	21
	KIDNEY, URETER & MAJ BLDR PROC FOR NON-NEOPL AGE >=70 &/OR C.C.	2.1561	0.6	7.2	-	72
	00	1.3649	5.8	4.4	-	20
	PROSTATECTOMY AGE >=70 AND/OR C.C.	2.0958	12.4	7.5	-	54
	PROSTATECTOMY AGE <70 W/O C.C.	1.0308	0.9	4.8	-	16
	MINOR BLADDER PROCEDURES AGE >=70 AND/OR C.C.	1.4545	7.1	4.8	-	21
	MINOR BLADDER PROCEDURES AGE <70 M/O C.C.	1.1876	5.9	4.2	•	12
	TRANSURETHRAL PROCEDURES AGE >=70 AND/OR C.C.	1.0370	4.4	3,3	-	14
	TRANSURETHRAL PROCEDURES AGE <70 W/O C.C.	0.7452	3.3	2.7	-	0
	URETHRAL PROCEDURES, AGE >=70 AND/OR C.C.	0.7844	4.4	3.6	-	16
	URETHRAL PROCEDURES, AGE 18-69 W/O C.C.	0.6461	3.5	2.4	-	11
	URETHRAL PROCEDURES, AGE 0-17	0.5325	2.8	2.2	-	80
	OTHER KIDNEY & URINARY TRACT O.R. PROCEDURES	2.6432	10.7	7.3	-	72
	RENAL FAILURE	1.9126	8.4	5.9	-	22
	ADMIT FOR RENAL DIALYSIS	42067.0	3.4	2.3	-	10
	KIDNEY & URINARY TRACT NEOPLASMS AGE >=70 AND/OR C.C.	1.3748	8.1	5.2	-	22
319 KIDN	KIDNEY & URINARY TRACT NEOPLASHS AGE <70 W/O C.C.	0.6935	4.3	3.1	-	15

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DRG MIMBER DESCRIPTION	CHAMPUS	ARITHMETIC MEAN 100	GEOMETRIC	CHAMPUS ARITHMETIC GEOMETRIC SHORT STAY LONG STAY	LONG STAY
	acres acres	ייבאו רוס	חבאו רטי		ועאבפעמרת
320 KIDNEY & URINARY TRACT INFECTIONS AGE >=70 AND/OR C.C.	0.9528	5.9	5.0	-	15
321 KIDNEY & URINARY TRACT INFECTIONS AGE 18-69 W/O C.C.	0.6524	4.4	3.7	-	1
322 KIDNEY & URINARY TRACT INFECTIONS AGE 0-17	0.5497	4.3	3.6	-	11
323 URINARY STONES AGE >=70 AND/OR CC, &/OR ESW LITHOTRIPSY	0.9396	3.6	2.6	-	11
324 URINARY STONES AGE <70 W/O C.C.	0.5417	2.4	1.9	•	9
325 KIDNEY & URINARY TRACT SIGNS & SYMPTOMS AGE >=70 AND/OR C.C.	0.8738	5.4	0.4	-	19
326 KIDNEY & URINARY TRACT SIGNS & SYMPTOMS AGE 18-69 W/O C.C.	0.6338	3.6	2.7	-	11
327 KIDNEY & URINARY TRACT SIGNS & SYMPTOMS AGE 0-17	0.5118	3.8	2.9	-	-
328 URETHRAL STRICTURE AGE >=70 AND/OR C.C.	0.6042	3.7	3.0	-	-1
329 URETHRAL STRICTURE AGE 18-69 W/O C.C.	0.5086	3.3	2.4	-	10
330 URETHRAL STRICTURE AGE 0-17	0.2788*	0.0	1.6	-	5
331 OTHER KIDNEY & URINARY TRACT DIAGNOSES AGE >=70 AND/OR C.C.	1.0961	6.8	4.6	-	12
332 OTHER KIDNEY & URINARY TRACT DIAGNOSES AGE 18-69 M/O C.C.	0.6032	3.7	2.8	-	11
333 OTHER KIDNEY & URINARY TRACT DIAGNOSES AGE 0-17	0.5760	3.8	2.6	-	12
334 MAJOR MALE PELVIC PROCEDURES WITH C.C.	2.0361	11.1	10.1	7	77
335 MAJOR MALE PELVIC PROCEDURES W/O C.C.	1.7552	8.5	8.0	7	15
336 TRANSURETHRAL PROSTATECTOMY AGE >=70 AND/OR C.C.	0.9604	5.1	4.6	2	10
337 TRANSURETHRAL PROSTATECTOMY AGE <70 W/O C.C.	0.7762	4.3	4.1	2	1
338 TESTES PROCEDURES, FOR MALIGNANCY	0.7460	3.8	2.7	-	13
339 TESTES PROCEDURES, NON-MALIGNANT AGE >=18	0.5773	2.5	1.9	-	1
340 TESTES PROCEDURES, MON-MALIGNANT AGE 0-17	0.4925	1.9	1.6	-	,
341 PENIS PROCEDURES	0.9421	4.1	3.2	-	13
342 CIRCUMCISION AGE >=18	0.4662	1.7	1.4	-	7
343 CIRCUMCISION AGE 0-17	0.6514	6.4	2.2	-	19
344 OTHER MALE REPRODUCTIVE SYSTEM O.R. PROCEDURES FOR MALIGNANCY	1.0783	8.4	3.9	-	15
345 OTHER MALE REPRODUCTIVE SYSTEM O.R. PROC EXCEPT FOR MALIG	0.8772	5.0	3.5	-	19
346 MALIGNANCY, MALE REPRODUCTIVE SYSTEM, AGE >=70 AND/OR C.C.	1.3703	9.3	5.0	-	22
347 MALIGNANCY, MALE REPRODUCTIVE SYSTEM, AGE <70 W/O C.C.	0.6337	4.1	3.0	-	13
348 BENIGN PROSTATIC HYPERTROPHY AGE >=70 AND/OR C.C.	0.6458	3.8	2.8	-	13

DRG	CHAMPUS A	RITHMETIC	GEOMETRIC	CHAMPUS ARITHMETIC GEOMETRIC SHORT STAY LONG STAY	LONG STAY
NUMBER DESCRIPTION	WEIGHT	MEAN LOS	MEAN LOS	THRESHOLD	THRESHOLD THRESHOLD
U	0.5316	2.4	2.0		
350 INFLAMMATION OF THE MALE REPRODUCTIVE SYSTEM	0.6710	4.5	3.8	-	1
	0.3333*	1.9	1.6	-	5
	0.5645	3.6	2.5	-	12
	1.9489	9.1	8.4	3	17
	1.5556	7.7	6.9	2	16
UTERINE, ADNEKA P	1.0703	5.7	5.3	2	10
	0.9374	5.6	5.0		12
UTERUS & ADENEXA	2.0570	0.6	8.0		20
	1.3064	4.9	5.9	2	12
UTERINE AND ADNEX	0.9826	5.1	8.4	2	•
	0.6131	2.9	2.3	-	60
	7689.0	3.0	2.3	-	
	0.3483	1.4	1.2		
	0.6366	3.2	2.6		10
	0.4863	2.1	1.7	-	
OTHER FEMALE REPR	1.2070	0.9	5.1	-	16
MALIGNANCY, FEMAL	1.1986	7.5	5.0	-	22
MALIGNANCY, FEMAI	0.6089	3.3	2.2	-	11
	0.6357	0.4	3.5	-	0
	0.4241	2.8	2.2	-	1
CROAKRAN SECTION	1.0878	0.9	5.4	2	11
CESAKEAN SECTION	0.9012	8.4	9.4	2	80
VAGINAL DELIVERY	0.8102	4.4	3.6	-	-=
	9997.0	2.7	2.4	-	5
VAGINAL DELIVERY	0.6730	3.0	2.8	-	2
372 ANGINAL DELIVERY WITH O.R. PROC EXCEPT STERIL AND/OR DAC	0.6817*	0.0	4.4	-	15
ATY DOCTOROUM AND PUSINGENETICM DIAGNOSES W/O O.R. PROCEDURE	0.4692	3.1	2.5	1	60
SIT PUSIFIER LOW AND POSTABORTION DIAGNOSES WITH O.R. PROCEDURE	0.6218	2.5	1.9	-	9

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DRG NUMBER DESCRIPTION	CHAMPUS	CHAMPUS ARITHMETIC GEOMETRIC SHORT STAY LONG STAY WEIGHT MEAN LOS THRESHOLD THRESHOLD	GEOMETRIC MEAN LOS	SHORT STAY THRESHOLD	THRESHOLD THRESHOLD
378 ECTOPIC PREGNANCY	0.8096	3.9	3.7	-	7
379 THREATENED ABORTION	0.3214	2.6	2.0	-	1
380 ABORTION W/O D&C	0.3211	1.8	1.5	-	4
381 ABORTION WITH D&C, ASPIRATION CURETTAGE, OR HYSTEROTOMY	0.3652	1.4	1.2	-	2
382 FALSE LABOR	0.1479	1.3	1.2	•	2
383 OTHER ANTEPARTUM DIAGNOSES WITH MEDICAL COMPLICATIONS	0.3560	3.2	2.6	-	6
384 OTHER ANTEPARTUM DIAGNOSES W/O MEDICAL COMPLICATIONS	0.3615	5.9	2.0	-	60
385 NEONATES, DIED OR TRANSFERRED TO ANOTHER ACUTE CARE FACILITY	0.8322	5.1	2.9	•	18
386 EXTREME IMMATURITY OR RESPIRATORY DISTRESS SYNDROME, NEOWATE	3.3194	16.8	10.6	-	27
387 PREMATURITY WITH MAJOR PROBLEMS	1.6986	11.9	8.5	-	25
388 PREMATURITY M/O MAJOR PROBLEMS	0.7994	7.6	5.3	-	22
389 FULL TERM NEONATE WITH MAJOR PROBLEMS	0.5024	9.4	3.7	-	12
390 NEONATES WITH OTHER SIGNIFICANT PROBLEMS	0.2257	3.3	2.9	-	80
391 NORMAL NEWBORNS	0.1390	2.7	2.4	-	9
392 SPLENECTOMY AGE >=18	3.0019	10.3	8.5	2	25
393 SPLENECTOMY AGE 0-17	1.8160	8.4	7.6	3	16
394 OTHER O.R. PROCEDURES OF THE BLOOD & BLOOD FORMING ORGANS	1.1131	5.3	3.2	-	20
395 RED BLOOD CELL DISORDERS AGE >=18	0.9470	5.5	4.0		19
396 RED BLOOD CELL DISORDERS AGE 0-17	0.6575	4.3	3.2	-	14
397 COAGULATION DISORDERS	0.7945	4.5	3.3	-	5
398 RETICULDENDOTHELIAL & IMMUNITY DISORDERS AGE >=70 AND/OR C.C.	1.3348	6.7	5.3	-	20
399 RETICULOENDOTHELIAL & IMMUNITY DISORDERS AGE <70 W/O C.C.	0.7116	8.4	3.6	-	16
400 LYMPHOMA OR LEUKEMIA WITH MAJOR O.R. PROCEDURE	3.6784	11.3	8.6	2	52
401 LYMPHOMA AND NON-ACUTE LEUKENIA WITH O.R. PROC WITH C.C.	2.0607	7.6	0.9	-	23
402 LYMPHOMA AND NON-ACUTE LEUKENIA WITH O.R. PROCEDURE W/O C.C.	1.2718	5.7	4.1	-	20
403 LYMPHOMA AND NON-ACUTE LEUKENIA WITH C.C.	1.7645	8.8	5.8	-	22
404 LYMPHOMA AND MON-ACUTE LEUKEMIA W/O C.C.	1.0364	5.7	3.8	-	20
405 ACUTE LEUKEMIA WITHOUT MAJOR O.R.PROCEDURE, AGE 0-17	1.6238	8.0	5.0	-	21
406 MYELOPROLIF DISORD OR POORLY DIFF NEOPLASM W MAJ O.R.PROC & C.C.	2.7904	12.4	9.5	2	97

DRG	CHAMPUS	CHAMPUS ARITHMETIC GEOMETRIC SHORT STAY LONG STAY	GEOMETRIC	SHORT STAY	LONG STAY	
NUMBER DESCRIPTION	WEIGHT	MEAN LOS	MEAN LOS		THRESHOLD THRESHOLD	32.44
407 MYELOPROLIF DISORD OR POORLY DIFF NEOPL W MAJ G.R.PROC W/O C.C.	1.3771	6.9	5.7	-	22	-
408 MYELOPROLIF DISORD OR POORLY DIFF NEOPL WITH OTHER O.R. PROC	1.0734	5.1	3.6	-	15	
409 RADIOTHERAPY	0.8893	6.2	4.3	-	12	
410 CHEMOTHERAPY	0.6178	5.9	2.3	•	80	
411 HISTORY OF MALIGNANCY M/O ENDOSCOPY	1.0036	7.0	3.0	•	20	
412 HISTORY OF MALIGNANCY WITH ENDOSCOPY	0.3388*	1.2.4	1.9		9	
413 OTHR MYELOPROLIF DISORD OR POORLY DIFF NEOPL DX AGE>=70 &/OR C.C	1.2413	8.2	5.1	-	22	1
414 OTHR MYELOPROLIF DISORD OR POORLY DIFF NEOPL DX AGE-70 W/O C.C.	7906.0	9.9	3.7	- Comment	20	-
415 O.R. PROCEDURE FOR INFECTIOUS & PARASITIC DISEASES	3.2853	13.4	9.3	-	92	
416 SEPTECEMIA AGE >=18	1.9711	0.6	6.8	-	23	
417 SEPTECEMIA AGE 0-17	0.7178	7.5	4.2	-	15	
418 POSTOPERATIVE & POST-TRAUMATIC INFECTIONS	1.1579	7.4	5.4	-	22	4
419 FEVER OF UNKNOWN ORIGIN AGE >=70 AND/OR C.C.	1.1611	6.5	5.5	-	17	
420 FEVER OF UNKNOWN ORIGIN AGE 18-69 W/O C.C.	0.9760	0.9	4.7	-	18	14-
421 VIRAL ILLNESS AGE >=18	0.5359	3.6	3.0		0	
422 VIRAL ILLNESS & FEVER OF UNKNOWN ORIGIN AGE 0-17	0.4118	3.3	2.8	-	00	
423 OTHER INFECTIOUS & PARASITIC DISEASES DIAGNOSES	1.0385	4.9	4.2	-	12	
424 O.R. PROCEDURES WITH PRINCIPAL DIAGNOSIS OF MENTAL ILLNESS	- 1		•			
425 ACUTE ADJUST REACT & DISTURBANCES OF PSYCHOSOCIAL DYSFUNCTION	1				•	
426 DEPRESSIVE NEUROSES						
427 NEUROSES EXCEPT DEPRESSIVE					-	
428 DISORDERS OF PERSONALITY & IMPULSE CONTROL				•		
429 ORGANIC DISTURBANCES & MENTAL RETARDATION			•		•	
430 PSYCHOSES			•	•		
431 CHILDHOOD MENTAL DISORDERS						
432 OTHER DIAGNOSES OF MENTAL DISORDERS				•	•	
433 ALCHOHOL/DRUG USE AND INDUCED ORGANIC MENTAL DISORDERS, LEFT AMA				•	•	
434 ALC/DRUG ABUSE, INTOX INDUCD MNTL SYN EXC DEPEND &/OR OTH SYMPT T				•	•	
435 ALCOHOL/DRUG DEPEND, DETOX AND/OR OTH SYMPTOMATIC TREATMENT			•		•	

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DRG NUMBER DESCRIPTION	CHAMPUS	CHAMPUS ARITHMETIC GEOMETRIC SHORT STAY LOWG STAY WEIGHT MEAN LOS THRESHOLD THRESHOLD	GEOMETRIC MEAN LOS	SHORT STAY LOWG STAY THRESHOLD THRESHOLD	LONG STAY THRESHOLD
436 ALCOHOL/DRUG DEPENDENCE WITH REHABILITATION THERAPY					
437 ALCOHOL/DRUG DEPENDENCE, COMBINED REHAB AND DETOX THERAPY					
438 NO LONGER VALID			•		400
439 SKIN GRAFTS FOR INJURIES	1.4983	7.9	4.7		21
440 WOUND DEBRIDEMENTS FOR INJURIES	1.5195	8.2	5.2		22
441 MAND PROCEDURES FOR INJURIES	0.9523	3.4	2.3	-	10
442 OTHER O.R. PROCEDURES FOR INJURIES AGE >=70 AND/OR C.C.	2.5787	9.5	6.3	-	23
443 OTHER O.R. PROCEDURES FOR INJURIES AGE <70 M/O C.C.	1.3343	5.8	3.5		20
444 MULTIPLE TRAUMA AGE >=70 AND/OR C.C.	0.9972	5.4	4.2	-	11
445 MULTIPLE TRAUMA AGE 18-69 W/O C.C.	0.8053	4.5	2.9	-	15
446 MULTIPLE TRAUMA AGE 0-17	0.6391	4.6	2.6	-	16
447 ALLERGIC REACTIONS AGE >=18	0.4913	2.9	2.2	-	80
448 ALLERGIC REACTIONS AGE 0-17	0.2952	2.3	1.8	-	9
449 POISONING AND TOXIC EFFECTS OF DRUGS AGE >=70 AND/OR C.C.	0.9045	5.4	3.3	-	20
450 POISONING AND TOXIC EFFECTS OF DRUGS AGE 18-69 W/O C.C.	0.4673	2.9	2.0	-	63
451 POISONING AND TOXIC EFFECTS OF DRUGS AGE 0-17	0.3600	2.3	1.7	-	9
452 COMPLICATIONS OF TREATMENT AGE >=70 AND/OR C.C.	1.0734	6.5	4.2	-	21
453 COMPLICATIONS OF TREATMENT AGE <70 W/O C.C.	0.4458	3.2	2.2	-	10
454 OTHER INJURIES, POISONINGS & TOXIC EFF DIAG AGE >=70 AND/OR C.C.	1,3881	4.1	2.6	-	14
455 OTHER INJURIES, POISONINGS & TOXIC EFF DIAG AGE <70 W/O C.C.	0.4340	2.0	1.6	-	5
456 BURNS, TRANSFERRED TO ANOTHER ACUTE CARE FACILITY	0.9577	8.2	5.3	-	22
457 EXTENSIVE BURNS W/O O.R. PROCEDURE	3.2280*	9.5	0.4		21
458 HOM-EXTENSIVE BURNS WITH SKIN GRAFTS	3.0929	15.0	10.6	-	27
459 NON-EXTENSIVE BURNS WITH WOUND DEBRIDEMENT & OTHER O.R. PROC	1.7401	7.8	6.2	-	23
460 NON-EXTENSIVE BURNS W/O O.R. PROCEDURE	0.8706	5.9	3.7	-	20
461 O.R. PROC WITH DIAGNOSES OF OTHER CONTACT WITH HEALTH SERVICES	0.9335	4.6	2.8	-	16
462 REHABILITATION	3.1150	25.4	18.7	2	35
463 SIGNS & SYMPTONS WITH C.C.	0.9359	4.9	4.7	-	12
464 SIGNS & SYMPTOMS W/O C.C.	0.6013	4.5	3.4	-	14

DRG NUMBER DESCRIPTION	CHAMPUS	ARITHMETIC MEAN LOS	GEOMETRIC MEAN LOS	CHAMPUS ARITHMETIC GEOMETRIC SHORT STAY LONG STAY WEIGHT MEAN LOS THRESHOLD THRESHOLD	LONG STAY THRESHOLD
465 AFTERCARE WITH HISTORY OF MALIGNANCY AS SECONDARY DX	0.3885	1.6	1.4	-	3
	0.4627	2.8	1.9	-	80
467 OTHER FACTORS INFLUENCING MEALTH STATUS	0.3429	2.7	2.0	•	7
468 UNRELATED OR PROC	1.6778	7.5	4.7	-	21
469 PDX INVALID AS DISCHARGE DIAGNOSIS			***************************************		The state of the s
470 UNGROUPABLE		•			
471 BILATERAL OR MULTIPLE MAJOR JOINT PROCEDURES OF THE LOWER EXTREM	4.3939	10.5		2	56
472 EXTENSIVE BURNS WITH O.R. PROCEDURE	12.3234*				0,
473 ACUTE LEUKEMIA W/O MAJOR O.R. PROCEDURE AGE > 17	4.7101	17.4	10.2		17

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Table 2—National Urban and Rural Adjusted Standardized Amounts, Labor/ Nonlabor, Cost-Share Per Diem, and Area Wage Indexes

[Editorial Note: This table will not appear in the Code of Federal Regulations]

The following standardized amounts are approximations. Revised amounts are being developed and the actual amounts will be published in the Federal Register in about one week.

National urban adjusted standard- ized amount	\$2,835.94
Labor portion	2,061.16 774.78
National rural adjusted standard- ized amount	2,518.41
Labor portion	1,942.45 575.96
Cost-share per diem for benefici- aries other than dependents of active duty members	175.00

Area Wage Indexes

The area wage indexes used under the CHAMPUS DRG-based payment system are those used under the Medicare PPS

as published in the Federal Register on June 10, 1987 (52 FR 22135).

Addendum 1—Health Program Benefit Agreement

[Editorial Note: This addendum will not appear in the Code of Federal Regulations.]

In order to receive payment under the Civilian Health and Medical Program of the Uniformed Services (CHAMPUS).

ba — as the provider of services agrees:

(a) To accept as payment for inpatient services provided to eligible beneficiaries, the CHAMPUS-determined allowable amount. This amount will be determined in accordance with the requirements of DoD 6010.8–R as published in the Federal Register on (insert date of publication).

(b) To refrain from billing the CHAMPUSeligible beneficiary for amounts which exceed the CHAMPUS-determined allowable amount except for services not covered by CHAMPUS as described in DoD 6010.8-R and for amounts which constitute the CHAMPUS beneficiary's liability for cost-share and deductible.

OCHAMPUS agrees:

(a) To pay the hospital the full allowable amount less any applicable cost-share and deductible amounts.

This agreement shall be binding on the provider and OCHAMPUS upon submission

by the provider of acceptable assurance of compliance with Title VI of the Civil Rights Act of 1964, section 504 of the Rehabilitation Act of 1973 as amended, and upon acceptance by the Director, OCHAMPUS, or his designee.

This agreement shall be effective until terminated by either party. The effective date shall be the date the agreement is signed by OCHAMPUS.

The agreement may be terminated by either party by giving the other party written notice of termination. Such notice of termination is to be received by the other party no later than 30 days prior to the date of termination. In the event of transfer of ownership, this agreement is terminated.

FOR PROVIDER OF SERVICES BY

FOR PROVIDER OF SERVICES B	13
Name	
Title	
Date FOR OCHAMPUS BY:	
Name	
Title	
Date	451

[FR Doc. 87-19684 Filed 8-31-87; 8:45 am]





Tuesday September 1, 1987

Part III

Department of Health and Human Services

Health Care Financing Administration

42 CFR Parts 405, 412, 413, and 466
Medicare Program; Changes to the
Inpatient Hospital Prospective Payment
System and Fiscal Year 1988 Rates; Final
Rule

Medicare Program: Changes to the DRG Classification System; Final Notice



DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

42 CFR Parts 405, 412, 413, and 466

[BERC-400-F]

Medicare Program; Changes to the Inpatient Hospital Prospective Payment System and Fiscal Year 1988 Rates

AGENCY: Health Care Financing Administration (HCFA), HHS. ACTION: Final rule.

SUMMARY: We are revising the Medicare inpatient hospital prospective payment system to implement necessary changes arising from legislation and our continuing experience with the system. One of these changes is the inclusion in the prospective payment system of hospitals located in Puerto Rico.

In addition, in the addendum to this rule, we describe changes in the methods, amounts, and factors necessary to determine prospective payment rates for Medicare inpatient hospital services. In general, these changes are applicable to discharges occurring on or after October 1, 1987. We also set forth the rate-of-increase limits for hospitals and hospital units excluded from the prospective payment system.

effective DATE: This final rule is effective on October 1, 1987. We refer the reader to section VII.A. of this preamble for a discussion of specific provisions that apply to specific periods.

FOR FURTHER INFORMATION CONTACT: Linda Magno, (301) 594-9343.

SUPPLEMENTARY INFORMATION:

I. Background

A. Summary

Under section 1886(d) of the Social Security Act (the Act), enacted by the Social Security Amendments of 1983 (Pub. L. 98-21) on April 20, 1983, a system for payment of inpatient hospital services under Medicare Part A (Hospital Insurance) based on prospectively-set rates was established effective with hospital cost reporting periods beginning on or after October 1, 1983. Under this system, Medicare payment is made at a predetermined, specific rate for each hospital discharge. All discharges are classified according to a list of diagnosis-related groups (DRGs). The regulations governing the inpatient hospital prospective payment system are located in 42 CFR Part 412.

Sections 1886(d)(1) (A), (C), and (D) of the Act provide for the implementation of the prospective payment system over a four-year transition period. During the transition period, payment to hospitals is based on a combination of the Federal prospective payment rates and hospital-specific rates, the proportions of which change with the hospital's cost reporting period. In addition, during that period, the Federal rate is a combination of regional and national rates, the proportions of which change with the Federal fiscal year.

B. Summary of the Provisions of the June 10, 1987 Proposed Rule

On June 10, 1987, we published a proposed rule in the Federal Register (52 FR 22080) to further amend the prospective payment system, as follows:

• We proposed to restructure the alcohol and drug abuse DRGs. We also proposed to reorder the surgical hierarchies in several Major Diagnostic Categories (MDCs). In addition, as required by section 1886(d)(4)(C) of the Act, as amended by section 9302(e)(1) of the Omnibus Budget Reconciliation Act of 1986 (Pub. L. 99–509), we proposed to adjust the DRG weighting factors for discharges in FY 1988.

 We proposed to change the methodology we use for computing the national average hourly wage that serves as the basis for indexing the area wage levels. We also proposed to update the wage index based on more

recent wage data.

• Under the authority of section 1886(d)(9)(A) of the Act, which was added by section 9304(a) of Pub. L. 99–509, we proposed that inpatient hospital services furnished by hospitals located in Puerto Rico are to be paid for under the prospective payment system beginning with discharges on or after October 1, 1987.

 We discussed several current provisions of the regulations in 42 CFR Parts 405, 412, 413, and 466 and set forth certain proposed changes concerning—

-Review of DRG assignments:

- —An increase in the prospective payment rates and rate-of-increase limits;
- -Payment for outlier cases;
- Payments to sole community hospitals;
- Referral center criteria and basis of payment; and
- Payment for services of nonphysician anesthetists.
- In the addendum to the proposed rule, we set forth proposed changes to the methods, amounts, and factors for determining the FY 1988 prospective payment rates. We also proposed new target rate percentages for determining the rate-of-increase limits for FY 1988

for hospitals and hospital units excluded from the prospective payment system.

In addition, the proposed rule discussed in detail the April 1, 1987 recommendations made by the Prospective Payment Assessment Commission (ProPAC). ProPAC is directed by section 1886(d)(4)(D) of the Act to make recommendations to the Secretary with respect to adjustments to the DRG classification and weighting factors and to report to Congress with respect to its evaluation of any adjustments made by the Secretary.

ProPAC is also directed, by the provisions of sections 1886(e)(2) and (e)(3) of the Act, to make recommendations to the Secretary on the appropriate percentage change factor to be used in updating the average standardized amounts beginning with FY 1986 and thereafter. We printed ProPAC's report, which includes its recommendations, as Appendix C to the proposed rule (52 FR 22167).

C. Number and Types of Public Comments

A total of 204 letters containing comments on the proposed regulations were received timely. More than half of the letters we received were protesting the end of the exclusion for alcohol/drug hospitals and units and the revised DRGs for alcohol and drug cases. Of the remaining letters, the only subject that was addressed by a majority of the commenters was the changes we proposed to make to the rural referral center regulations.

The contents of the proposed rule, the public comments, and our responses to the comments are discussed through this document in the appropriate sections. As discussed below in section II of this preamble, the comments concerning the restructuring of the alcohol and drug abuse DRGs and the changes made to the surgical hierarchies are addressed in a separate notice concerning changes to the DRG classification system published elsewhere in this issue of the Federal Register.

In addition, we note that, in this document, we are not responding to comments that raised issues not specific to the proposals we made. These issues include the criteria for receiving periodic interim payments, the general prospective payment methodology (which is, for the most part, set by law), and the level of care permitted to be furnished in inpatient hospital beds.

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There is one general comment that we are responding to here rather than in one of the more issue-specific areas below.

Comment: A number of commenters, particularly major associations and organizations, stated that we did not make available to the public sufficient information and documentation that would provide the hospital industry a basis on which to respond appropriately to the provisions of the proposed rule.

Response: As we have stated in previous final rules on the prospective payment system in response to similar comments (see 50 FR 35657 and 51 FR 31491), all of the disclosable data files used in computing the prospective payment rates, DRG weighting factors, and impact analyses are available to the public upon request under the Freedom of Information Act (5 U.S.C. 552). In fact, many institutions and organizations have requested and received these files. If we were to publish data from these files in the Federal Register, along with the detailed computations used in deriving factors such as the DRG weights, the proposed notice and comment procedure would be unnecessarily complicated because many potential commenters would be inundated with material that might be of little or no interest to them.

However, each year, while we are developing the policies and changes in current policies that we will be presenting in the next proposed rule, we meet with interested parties when necessary. Also, ProPAC has open meeting about many of the same issues that concern us. Thus, the hospital industry, through attendance at these meetings, can become knowledgeable about the issues that we are examining.

We believe that these opportunities, coupled with the statutory requirement for the 60-day public comment period (section 1871(b)(1) of the Act), provide the hospital industry with ample opportunity to obtain sufficient data and information on which to base their comments.

II. Changes to DRG Classifications and Weighting Factors

A. Background

Under the prospective payment system, we pay for inpatient hospital services on the basis of a rate per discharge that varies according to the DRG to which a beneficiary's stay is assigned. The formula used to calculate payment for a specific case takes an individual hospital's average payment rate per case and multiplies it by the weight of the DRG to which the case is assigned. Each DRG weight represents the average resources required to care for a case in that particular DRG relative to the national average of resources consumed per case. Thus, cases in a

DRG with a weight of 2.0 would, on average, require twice as many resources as the average case.

Congress recognized that it would be necessary to recalculate the DRG relative weights periodically to account for changes in resource consumption. Accordingly, section 1886(d)(4)(C) of the Act, as originally added to the Act by Pub. L. 98-21, required that the Secretary adjust the DRG classifications and weighting factors effective for discharges occurring in FY 1986 and at least every four fiscal years thereafter. These adjustments were to be made to reflect changes in resource consumption, treatment patterns, technology, and any other factors that may change the relative use of hospital resources.

Section 9302(e) of Pub. L. 99-509 revised section 1886(d)(4)(C) of the Act to require that we adjust the DRG classifications and weighting factors annually beginning with discharges occurring in FY 1988. The majority of the proposed changes to the DRG classification system for discharges occurring in FY 1988 were discussed in a notice published in the Federal Register on May 19, 1987 (52 FR 18877). However, as a part of the proposed rule, we addressed two of the reclassification issues; that is, the alcohol and drug abuse DRGs and surgical hierarchies. The comments we received on the restructuring of the alcohol and drug abuse DRGs and the revisions to the surgical hierarchies as well as any other changes we are making to the DRG classification system are discussed in a separate notice published elsewhere in this issue of the Federal Register. However, in section II.B, below, we discuss the comments received on the alcohol and drug abuse DRGs that conern issues other than the restructuring.

We are recalibrating the DRG weights as discussed below. In addition, we have revised § 412.60(d), which describes how often we revise the DRG classification and weighting factors, so that it conforms to the law as amended by Pub. L. 99–509.

Recalibration of DRG Weights

One of the basic issues in recalibration is the choice of a data base that allows us to construct relative DRG weights that most accurately reflect current relative resource use. The previous recalibration of DRG weights, which was published as a part of the FY 1986 prospective payment final rule, used hospital charge information from the central office enrollment file and the FY 1984 Part A Tape Bill (PATBILL) data set to create the MEDPAR file. For a discussion of the options we considered

and the reasons why we chose to use charge data for the FY 1986 recalibration, we refer the reader to the June 10, 1985 proposed rule (50 FR 24372) and the September 3, 1985 final rule (50 FR 35652).

We proposed to use the same methodology for the FY 1988 recalibration as we did for FY 1986. That is, we recalibrated the weights based on charge data for Medicare discharges occurring in FY 1986. However, we used the FY 1986 Medicare provider analysis and review (MEDPAR) file rather than the PATBILL data used in the DRG recalibration that was effective for discharges occurring in FY 1986. The MEDPAR file contains the same data as the PATBILL file but is in a simplified. reformatted record layout. MEDPAR is now based on fully-coded diagnostic and surgical procedure data for all Medicare inpatient hospital bills rather than for a 20-percent sample of beneficiaries as was the case in the FY 1986 recalibration. In addition, because the DRG weights are to be used to calculate prospective payments to hospitals in Puerto Rico beginning with discharges on or after October 1, 1987 and to alcohol/drug hospitals and units effective with cost reporting periods beginning on or after October 1, 1987 bills from these hospitals were included in the data set used to recalibrate the weights.

The proposed recalibrated DRG relative weights were constructed from FY 1986 MEDPAR data received by HCFA through February 1987 and were based on almost 90 percent of all Medicare discharges occurring in FY 1986 from those hospitals that will be subject to the prospective payment system in FY 1988. The MEDPAR file data included approximately 9.4 million Medicare discharges (erroneously indicated as 9.5 million in the proposed rule). The MEDPAR file thorugh June 1987 includes 9.7 million or more than 90 percent of FY 1986 discharges, and this is the file used to calculate the weights set forth in Table 5 of this final rule.

The methodology used to calculate the DRG weights from the MEDPAR file is as follows:

- All the claims were regrouped using the revised DRG classifications set forth in a notice published elsewhere in this issue of the Federal Register.
- Charges were standardized to remove the effects of differences in area wage levels, indirect medical education payments, disproportionate share payments and, for hospitals in Alaska and Hawaii, the applicable cost-of-living adjustment.

 The average standardized charge per DRG was calculated by summing the standardized charges for all cases in the DRG and dividing that amount by the number of cases classified in the DRG.

 We then eliminated statistical outliers using the same criterion as was used in computing the current weights. That is, all cases outside of 3.0 standard deviations from the mean of the log distribution of charges per case for each DRG were eliminated.

· The average charge for each DRG was then recomputed excluding the statistical outliers and divided by the national average standardized charge per case to determine the weighting factor.

· In establishing the weighting factor for heart transplants (DRG 103), we used data for the 46 heart transplant cases (from 20 hospitals) in the FY 1986 MEDPAR file consistently with the methodology for all other DRGs. After removing statistical outliers, there were 45 cases on which the weight was based. Because heart transplants were not a Medicare covered service in FY 1986, we verified that the 20 hospitals whose cases were used to establish the weight were in fact hospitals that perform heart transplants.

 No adjustments were made to the charges to remove capital-related and direct medical education costs, as hospitals do not make discrete charges for these components of inpatient

hospital services.

· Kidney acquisition costs continue to be paid on a reasonable cost basis but, unlike other excluded costs, kidney acquisition costs are concentrated in a single DRG (DRG 302, Kidney Transplantation). For this reason, it was necessary to make an adjustment to prevent the relative weight for DRG 302 from including the effect of kidney acquisition costs, since these costs are paid separately from the prospective payment rate. Kidney acquisition charges were subtracted from the total charges for each case in DRG 302 prior to computing the average charge for the DRG and prior to eliminating statistical outliers.

The weights developed according to the methodology described above, using the revised GROUPER program, result in an average case weight that is slightly different from the average case weight before recalibration. Therefore, the new weights were normalized by an adjustment factor so that the average case weight after recalibration is equal to the average case weight prior to recalibration. This adjustment is intended to ensure that recalibration by itself neither increases nor decreases

total payments under the prospective payment system.

When we recalibrated the DRG weights for FY 1986, we set a threshold of 10 cases as the minimum number of cases required to compute a reasonable weight. At that time, there were 30 DRGs that contained no cases or fewer than 10 cases. We proposed to use that same case threshold in recalibrating the DRG weights for FY 1988. In addition, in the FY 1986 recalibration, we computed the weight for the 30 low-volume DRGs by adjusting the original weights of these DRGs by the percent change in the weight of the average case in the remaining DRGs. We proposed to use this same methodology for the FY 1988 recalibration.

Using the FY 1986 MEDPAR data set, there were 32 DRGs that contain fewer than 10 cases. Since we have no new data upon which to base the weights for these DRGs, we proposed to hold their current weight constant. This preserves the relationship between the weighting factor for each low-volume DRG and the average case weight for all Medicare cases.

In accordance with our September 3, 1986 final rule concerning changes to the inpatient hospital prospective payment system and FY 1987 rates (51 FR 31454), the exclusion of alcohol/drug treatment hospitals and units was extended through cost reporting periods beginning before October 1, 1987. The extension was intended to permit completion of analyses of a record reabstraction study conducted by the Alcohol, Drug Abuse, and Mental Health Administration (ADAMHA) in concert with the National Institute of Mental Health, the National Institute on Drug Abuse, the National Institute on Alcohol Abuse and Alcoholism, the Office of the Assistant Secretary for Planning and Evaluation, and HCFA.

The record reabstraction study was designed to evaluate the current structure of the alcohol/drug DRGs (MDC 20) and to identify variables not currently included in the DRG logic for MDC 20, such as patient age, disability status, complications and comorbidities (CC) and polysubstance use, that might account for additional variation in patient resource use. Based on the analyses and recommendations of ADAMHA, and our own analyses of the FY 1985 and FY 1986 MEDPAR records for all Medicare discharges in MDC 20, we proposed to reconfigure the alcohol/ drug DRGs as described in the June 10, 1987 proposed rule.

We received over 100 comments regarding the proposed reconfiguration of the alcohol/drug DRGs, the need for an impact analysis estimating the effect of paying alcohol/drug hospitals and units under the prospective payment system, and the proposed recalibrated weights and outlier thresholds for the alcohol/drug DRGs. Comments on the proposed restructuring of the alcohol/ drug DRGs are addressed in the final notice of DRG classification changes published elsewhere in this issue of the Federal Register. The remaining comments are addressed here, along with the other comments we received on our proposed recalibration.

Comment: Several commenters urged that we refine the DRG system to reflect more accurately the severity of a patient's illness or condition within any given DRG. The commenters urged that we make every effort to comply with section 9305(a) of Pub. L. 99-509, which directs the Secretary, by October 1988, to develop a legislative proposal to improve the DRG system and, in particular, to account for variations in severity of illness and case complexity

as a part of that proposal.

Response: We are currently evaluating a number of different severity of illness measures. There is no agreement within either the government or the hospital industry as to which of these systems, if any, is the most appropriate for use in a national payment program such as Medicare. In the meantime, we are continuing to

develop our proposal. Comment: One commenter expressed concern about the proposed weighting factor assigned to DRG 103 (heart transplant). Based on a study, the commenter maintained that the proposed weight (13.9614) is too low because it does not adequately account for the higher cost-to-charge ratio in heart transplant cases compared to average cost-to-charge ratios. That is, the commenter believes that the charges for heart transplant patients represent greater resource intensity (a higher costto-charge ratio) than is the case in charges for nontransplant patients. In this connection, another commenter suggested that the appropriateness of recalibrating the DRG weights using charge data alone should be reevaluated.

Response: As we stated in the proposed rule, the weighting factor for heart transplants was developed in a manner similar to that used to develop the weighting factors for all other DRGs. To the extent that the weighting factors for all DRGs are developed from FY 1986 charge information from Medicare patient bills, it is appropriate that the weighting factor for heart transplants be developed in the same way. Otherwise, the weighting factors will not be a

consistent measure of the relative resource use among DRGs. We note that the 55 heart transplant cases used to construct the final weighting factor for DRG 103 include many cases from several of the hospitals already approved to furnish Medicare-covered heart transplants, as well as those hospitals that we anticipate will receive approval in the near future.

We believe that the commenter's real concern is the same as the second commenter's, that the current method of determining the DRG weights based on charge data alone needs to be reviewed based on more recent data. We will be evaluating whether any changes may be appropriate with respect to recalibrating the DRG weighting factors. As indicated above, we are also evaluating a number of severity-of-illness adjustments to determine whether such an adjustment

should be adopted.

We note that when we originally adopted charge-based DRG weighting factors as a part of the FY 1986 recalibration, we did explore what effect, if any, the different cost-to-charge ratios across hospital departments would have on these weights. Our analysis demonstrated that, contrary to expectation, DRGs with high proportions of cost in "undercharged" services do not, in fact, have uniformly lower charge-based weights. Moreover, as we indicated in the September 3, 1985 final rule (50 FR 35655), the effects of a hospital subsidizing "undercharged" services with high-charge services are more complicated than might at first appear, and, in fact, we believe that the effects will, in many cases, offset one another.

Comment: One commenter observed that the proposed rule was silent with respect to the treatment of heart acquisition costs in the establishment of

the heart transplant weight.

Response: As indicated in the April 6, 1987 notice of HCFA ruling extending coverage to heart transplantations (52 FR 10935), heart acquisition costs will be paid separately, as a pass-through, for the time being. Accordingly, for the 55 heart transplant cases in the updated MEDPAR file used for recalibration, we subtracted from the total charges of each case an estimate of heart acquisition charges prior to computing the average charge for the DRG and prior to eliminating statistical outliers. This adjustment is identical to that used for removing kidney acquisition charges from cases in DRG 302 (Kidney Transplant).

Because current MEDPAR data do not separately identify heart acquisition charges, it was necessary to estimate such charges. Limited data available to

us from two hospitals that have already been approved as heart transplant centers revealed a range from \$2,500 at one facility to \$10,000 at another. One hospital in a State under waiver from the prospective payment system charges \$11,000 for heart acquisition. We also considered using the mean charge for kidney acquisition based on MEDPAR records for FY 1986, which was \$11,800. In light of limited data, we decided to use the mode charge for kidney acquisition in FY 1986, or \$7,000, as our best estimate of heart acquisition charges. We believe this figure is a reasonable estimate of charges associated with heart acquisition since it is within the range of the limited data available to us from three hospitals furnishing heart transplants under the Medicare program.

Comment: One commenter asked for reassurance that the renormalization of the DRG weights has been performed and that budget neutrality based on the changes to the DRGs has been

maintained.

Response: In renormalizing the DRG weights as we described above, we

maintained budget neutrality.

Comment: A number of commenters. predominantly alcohol/drug treatment facilities and the National Association of Addiction Treatment Providers (NAATP), expressed concern that the proposed weighting factors and lengths of stay for the alcohol/drug DRGs are based on the combined experience of both excluded alcohol/drug treatment facilities and short-stay hospitals furnishing services in "scatter beds," that is, in medical-surgical beds rather than in units organized exclusively around the provision of comprehensive alcohol/drug rehabilitative services. These commenters allege that the averaging effects work systematically to their disadvantage and that the resulting payments will force them to reduce services inappropriately to Medicare beneficiaries, especially in the DRGs for rehabilitation.

Response: The weighting factors for all DRGs are based on the average standardized charges of Medicare cases in each DRG relative to the average standardized charge for all Medicare cases in all DRGs. We use Medicare billing records from all hospitals and units subject to the prospective payment system under section 1886(d) of the Act. This policy has been in effect since the beginning of the Medicare prospective payment system. We have no reason to limit the Medicare cases used in recalibration to a subset of hospitals, particularly when coverage of all medically necessary items and services is not thus limited to that same subset of hospitals. We note that about 70 percent of all Medicare discharges in MDC 20 are from short-stay hospitals already subject to the prospective payment system.

As to the commenters' concerns that the lengths of stay published in Table 5 of the proposed notice are too low to reflect appropriate treatment patterns of patients receiving rehabilitation services, we reiterate that the length-ofstay figures published in the table of DRG relative weights are illustrative only and reflect historical lengths of stay for Medicare beneficiaries classified within each DRG. They represent neither treatment norms nor limitations on Medicare coverage or benefits. In addition, they do not represent expectations regarding future lengths of stay. In short, the lengths of stay are published for information purposes only and, except for the use of the geometric mean length of stay in calculating day outlier and transfer payments, have no bearing whatsoever on Medicare payment for inpatient hospital services.

For additional information on the distribution of Medicare lengths of stay by DRG, we refer the reader to Tables 7a and 7b in the Addendum to this final

rule.

Both tables display the lengths of stay by DRG for cases at the 10th, 25th, 50th, 75th and 90th percentiles of the distribution of all cases within that DRG. Table 7a shows the distribution of cases classified in accordance with the current DRG definitions while table 7b shows the distribution of cases regrouped in accordance with the DRG classifications that will be in effect for FY 1988. Both tables are based on the MEDPAR file of FY 1986 Medicare discharges from hospitals subject to the prospective payment system received in HCFA central office through June 1987. (For further information on the FY 1988 DRG classifications, see the final notice of changes to the DRG classification system, published elsewhere in this issue of the Federal Register.)

With respect to the comparative lengths of stay between short-stay hospitals and excluded alcohol/drug hospitals and units, we note that the Medicare average length of stay for discharges paid under the prospective payment system has declined by about 17 percent since the inception of that system. We believe that this drop contributes significantly to the differences that exist between the average lengths of stay in short-stay hospitals, which have now been paid subject to the prospective payment system for nearly four years, and

alcohol/drug hospitals and units that have been excluded from prospective payment system. We anticipate that once these excluded hospitals and units are brought into the prospective payment system, they would respond to its incentives, similarly to all other hospitals, by changing practice patterns, increasing productivity, and substituting lower-cost for higher-cost inputs.

In addition, many of the comments appear to reflect an unfounded assumption that all alcohol/drug cases paid under the prospective payment system represent treatment in scatter beds of hospitals that, in the words of one commenter, "lack the necessary program and staffing components to provide a comprehensive continuum of care." We believe the evidence does not support this conclusion. In point of fact, some hospitals certified and participating in Medicare as short-stay general hospitals identify themselves in the annual survey of the American Hospital Association as either alcohol specialty hospitals or general hospitals with organized alcoholism/chemical dependency inpatient units. Futhermore, under §§ 412.23 and 412.32, which specify requirements governing this exclusion, an alcohol/drug hospital or unit not excluded for its cost reporting period beginning during FY 1985 may not be excluded for subsequent cost reporting periods. Hence, facilities that first met the requirements as excluded alcohol/drug hospitals or units during their cost reporting period beginning on or after October 1, 1985, could not be excluded and would appear as shortstay hospitals in Medicare program files. Finally, we have found that of all alcohol/drug cases from short-stay hospitals, some 10 percent are cases billed under the prospective payment system from hospitals that have an excluded alcohol/drug unit.

If we are to assume that these hospitals do not deliberately treat some types of patients in their excluded units and other types of cases under the prospective payment system, and that they consistently furnish all necessary and appropriate care regardless of whether a patient is treated in an included or excluded bed, the data suggest that such hospitals have already made adjustments to respond to the incentives of the prospective payment system. The resource intensity of prospective payment cases from hospitals with excluded units, in terms of both standardized charges and mean length of stay, is more similar to that of other prospective payment cases in MDC 20 than to that of cases from excluded alcohol/drug units.

Comment: A few commenters observed that the proposed weighting factors are "biased" toward detoxification services compared to rehabilitation services.

Response: We believe that these commenters' concerns reflect a misunderstanding of the composition of DRGs 434 and 435, either as currently structured or as revised for FY 1988. These two DRGs include patients receiving detoxification services and/or symptomatic treatment. Detoxification therapy (code 94.25) is reported in fewer than 60 percent of the cases in these two DRGs. Other symptomatic treatment encompasses any other services furnished to a patient with a principle diagnosis in MDC 20 and may include medical management of other conditions aggravated by a patient's alcohol/drug abuse, such as dehydration and gastrointestinal bleeding, as well as occasional surgical intervention.

We believe that it is the range of services under the rubric "other symptomatic treatment" that contribute to a weighting factor that is higher than commenters appear to have expected. This inference is bolstered by our finding that presence or absence of detoxification therapy contributed insignificantly to explaining differences in resource use. Based on extensive review of these cases and evaluation of several alternative configurations, we concluded that the presence or absence of non-MDC 20 CCs was the strongest distinguishing characteristic among these patients. Further, in terms of the number of diagnoses and procedures reported on their Medicare bills, patients who would be assigned to DRG 434 as revised appear to be sicker and more complex to treat, on average, than most other patients in MDC 20.

Comment: A number of the commenters argued that the treatment practices of their specialized units and facilities are more comprehensive than the practices of prospective payment hospitals and that additional payments ought to be made to reflect this difference in treatment. In addition, it was suggested that treatments for alcohol and drug abuse patients provided by nonexcluded hospitals are inadequate and ought not to be covered under the Medicare program at all. The implicit assumption underlying these comments is that the average length of stay for rehabilitation cases in prospective payments hospitals is too short to constitute rehabilitation therapy as defined by Medicare program instructions.

Response: We do not agree with these commenters. Comments we have

received on this issue in response to previous proposed rules, most notably in the June 10, 1985 proposed rule, made it clear that there is considerable diversity of opinion in the field of alcohol and drug abuse treatment. As we noted in the final rule published on September 3, 1985 (50 FR 35651), we recognize that there are a variety of settings in which detoxification and rehabilitation can take place. We have not attempted to specify the explicit comments of such services or expected lengths of stav associated with particular modes of treatment nor do we believe that this would be appropriate, especially given the diversity of medical opinion in the field. Although commenters have tended to argue strongly in support of the efficacy of the treatment practices to which they adhere, we do not believe that any of them have presented evidence that such treatments are the optimum or preferred modes of treatment or, conversely, that other treatment modalities are conclusively ineffective and, therefore, should be deemed noncovered. Thus, we believe that the detoxification and rehabilitation services performed in nonexcluded prospective payment hospitals (nearly 70 percent of these services) must be given due weight in determining payment for the hospitals in question under the prospective payment system.

As to expectations of the commenters regarding appropriate lengths of stay for rehabilitation therapy based on their own facilities, we compared the average length of stay for cases in DRG 436 (Alcohol/Drug Dependence with Rehabilitation Therapy) from excluded alcohol/drug hospitals and hospitals already subject to the prospective payment system. In both FY 1985 and FY 1986, rehabilitation cases from hospitals under the prospective payment system have higher average lengths of stay than rehabilitation cases from excluded alcohol/drug hospitals, although both types of hospitals have shorter lengths of stay than rehabilitation cases from excluded alcohol/drug units. This finding holds regardless of whether cases are grouped in accordance with the current DRG 436 definition or with the revised structure for DRG 436. This comparison confirms our belief that there is as much diversity in treatment patterns and modalities among the excluded alcohol/drug hospitals and units as there is between prospective payment hospitals and excluded specialized facilites and that, in the face of such diversity, it is neither necessary nor appropriate to preserve the current payment distinction (the exclusion of alcohol/drug hospitals and units) or to

establish a new payment distinction within the prospective payment system

based on provider type.

Comment: At least one commenter referenced a study by the National Institute of Mental Health (NIMH), which analyzed psychiatric, alcohol, and drug Medicare inpatient admissions. The commenter expressed concern that the DRG recalibration does not take into account the fact "that all providers rendering care to individuals with these conditions are not similar with respect to the program of care provided," and there would be "a negative differential impact on those hospitals and distinct part units which specialize in the treatment of alcoholism and drug dependency cases." The NIMH study was quoted as documenting that "DRG payments to exempt hospitals and units would have in most cases been less than their historical costs for treating alcoholism and drug dependency admissions, while for the vast majority of non-exempt hospitals the DRG based payment would exceed historical costs." A second NIMH study was cited. addressing the differential in average costs for alcohol and drug discharges from specialty hospitals and general

Response: NIMH did conduct a study of Medicare alcohol, drug, and psychiatric inpatient admissions in four States analyzing the impact of a variety of patient classifications on payment to hospitals with and without excluded units. The finding quoted by the commenter refers to all of these admissions and, as such, is inappropriate for application to alcohol/ drug admissions only. The study did not distinguish from psychiatric cases; furthermore, the study used the original DRG constructions and weight and. therefore, cannot be construed to reflect the impact of the revised DRGs.

In addition to being invalid to extrapolate the findings of the study exclusively to alcohol/drug cases, the project looked at 1982 and 1983 data, at which time there was no distinction made between detoxification and rehabilitation protocols. For those hospitals under analysis, there was no distinction made between types of

treatment.

The second NIMH study cited by commenters is currently in progress; all findings are preliminary. This study also uses the original DRGs for alcohol/drug cases, and since the study data base consists of Medicare records from FY 1984 and FY 1985, a substantial number of cases predate the Medicare billing instructions that required explicit coding of detoxification and rehabilitation services for alcohol/drug cases.

Comment: Many commenters contended that the Medicare intermediary guidelines defining rehabilitation therapy and detoxification procedures need to be clarified and made "stringent enough to assure that only those providers capable of delivering said procedures in fact be reimbursed for doing so." Specifically, these commenters recommended that the guidelines incorporate into the definition of rehabilitation treatment the requirement that these services be furnished only in identifiable hospitals and units that meet the current criteria for exclusion as an alcohol/drug hospital and unit.

Response: With respect to the commenters' intimations that short-stay hospitals are not capable of furnishing alcohol/drug rehabilitation services and that the quality of such services is suspect, we note that during FYs 1984 through 1986, PROs reviewed more than 40 percent of all inpatient hospital stays through a combination of targeted reviews and random sampling. We have no evidence to suggest that there were problems in either the quality of services furnished to patients with alcohol/drug diagnoses or the validity of the DRG assigned based on review of the medical

record.

For the reasons stated in response to previous comments, we believe that it is inappropriate for the Medicare program to specify the types of settings, or the organizational structure thereof, in which alcohol/drug rehabilitation services may be furnished. To do so would either impose organizational and staffing requirements on a significant number of short-stay hospitals currently furnishing these services or severely limit beneficiary access to alcohol/drug rehabilitation services, 40 to 50 percent of which are furnished in short stay general hospitals. There is no conclusive evidence that such organizational and structural changes would produce improved treatment outcomes.

We believe the commenters are misinterpreting the ability of a facility to meet the exclusion criteria as assessments of the efficacy of the treatment they furnish, drawing the erroneous conclusion that certain services should then be covered only in facilities that meet said criteria. We note, however, that the exclusion criteria were tightly constructed in order to ensure that the alcohol/drug exclusion would not serve merely as a vehicle for an across-the-board exclusion of alcohol/drug cases from the prospective payment system.

When the prospective payment system was implemented, the clear purpose was to base payment for care

on the actual experience of most hospitals, and the DRG system included all the types of care these hospitals furnished. Only four types of specialty hospitals were specifically excluded by Congress from the prospective payment system: Rehabilitation hospitals and units, psychiatric hospitals and units, childrens' hospitals, and long-term care hospitals. These exclusions were put in the law as a result of a Congressional belief that these types of hospitals were sufficiently different from short-stay general hospitals as to warrant further study and separate treatment. In particular, the diagnoses and procedures associated with the patients of those institutions were perceived as inadequate to explain significant variations in patient resource utilization.

No such exclusion was granted for alcohol and drug hospitals and units (although such hospitals and units did and do have the option of qualifying under the psychiatric hospital and unit provision). We established the current exclusion relating to alcohol and drug abuse hospitals and units in recognition of the fact that the original MDC 20 DRGs failed to distinguish between detoxification and rehabilitation and the resource differences associated therewith. We have completed this task and, in addition to distinguishing between detoxification and rehabilitation, have made other modifications to the alcohol/drug DRGs. Accordingly, we believe there is no longer a basis or need for maintaining the alcohol/drug hospital and unit exclusion.

Comment: Virtually all commenters urged us to perform an impact analysis on the effect of paying currently excluded alcohol/drug hospitals and units under the prospective payment system.

Response: We refer readers to Appendix A of this final rule for the requested impact analysis.

III. Changes to the Hospital Wage Index Methodology

Section 1886(d)(2)(C)(ii) of the Act required, as a part of the process of developing separate urban and rural standardized amounts for FY 1984, that we standardize the average cost per case of each hospital for differences in area wage levels. Section 1886(d)(2)(H) of the Act requires that the standardized urban and rural amounts be adjusted for area variations in hospital wage levels as part of the methodology for determining prospective payments to hospitals. To fulfill both requirements, we constructed an index that reflects average hospital wages in each urban or

rural area relative to a national average

hospital wage.

For purposes of determining the prospective payments to hospitals in FY 1984 and FY 1985, we constructed the wage index using calendar year 1981 hospital wage and employment data obtained from the Bureau of Labor Statistics' ES 202 Employment, Wages and Contributions file for hospital workers. Subsequently, for FY 1986, the September 3, 1985 final rule set forth a revised hospital wage index that was based on an HCFA survey of 1982 hospital wage and salary data as well as data on paid hours in hospitals. That wage index was developed in an attempt to overcome the limitation of the BLS data with regard to full-time and part-time employment. As a result of the provisions of section 9103(a) of the Consolidated Omnibus Budget Reconciliation Act of 1985 (Pub. L. 99-272), application of the revised wage index was postponed from discharges occurring on or after October 1, 1985 to discharges occurring on or after May 1, 1986. The method used to compute the current HCFA wage index was set forth in detail in the September 3, 1985 final rule (50 FR 35661). In the September 3, 1986 final rule, we stated that we were collecting data as part of the audit of cost reports for the first year of the prospective payment system (FY 1984) in order to update the HCFA wage index (51 FR 31499).

Under the authority of section 9103(a) of Pub. L. 99–272, we proposed to make a change in the methodology for computing the national average hourly wage, which serves as the basis for indexing the area wage levels. We also proposed to adopt a blended wage index that incorporates the wage index based on 1982 data but computed using the proposed revised methodology discussed below and a new wage index based on 1984 data and also computed using the proposed methodology.

Currently, the wage index value for an area is computed by dividing the area's average hourly wage by the national average hourly wage. The national average hourly wage is computed by summing the average hourly wages for each area and dividing by the number of areas. Thus, the average hourly wage for each area is weighted equally in determining the national average hourly wage regardless of the number of hospitals or the size of the hospital labor force in the area.

Using the current methodology (that is, an area-weighted national average hourly wage) leads to a problem whenever the wage data for hospitals in an area are adjusted or when hospitals are reclassified from one area to

another. When either of these situations occurs, the national average hourly wage is affected, and thus the wage index values of all areas change.

Because of this problem, we proposed to compute the national average hourly wage by dividing the total wages for all hospitals by the total paid hours. This results in a wage index that is hourweighted rather than area-weighted. If the national average hourly wage is hour-weighted, there is minimal, if any, impact on that national average when the wage data for a particular area are adjusted.

While the proposed change in methodology for computing the national average wage does not affect the relative wage levels among areas. it does result in lower index values for all areas relative to the national average hourly wage, since the national average hourly wage is higher under our revised methodology than it would be if computed on an area-weighted basis. Therefore, in the addendum to the proposed rule (52 FR 22102), we proposed to restandardize the Federal payment amounts to reflect the proposed new method of computing the national average hourly wage.

In addition to proposing use of a revised methodology for computing the national average hourly wage, we also proposed, under the exceptions and adjustments authority in section 1886(d)(5)(C)(iii) of the Act, to adopt a blended wage index that incorporates both 1982 and 1984 wage data from prospective payment hospitals. The proposed index was based on area wage index values computed from 1982 data on an hour-weighted basis and area wage index values computed from 1984 data on an hour-weighted basis, equally weighted to produce average area wage index values.

The method used to compute the wage index is as follows:

Procedure I: Recomputation of the 1982 wage index on an hour-weighted basis.

Step 1—Each of the non-Federal acute care hospitals subject to the prospective payment system for which 1982 data were received was classified into its appropriate urban or rural area based on the current definitions of urban and rural areas used in the prospective payment system.

Step 2—For each hospital, the total gross hospital salaries were inflated from the end of the hospital's cost reporting year through the end of calendar year 1982, using the 1982 annual rate of increase in the wages and salaries portion of the hospital market basket. This was done to eliminate any

distortion caused by differing hospital cost reporting years.

Step 3—For each hospital, the inflated gross hospital salaries computed in step 2 were divided by the reported number of total paid hours to yield an average hourly wage. Hospitals with an aberrant hourly wage, which was defined as an hourly wage either less than \$3.35 (the minimum wage in 1982) or greater than \$19.58 (2.5 times the 1982 national average hourly hospital wage as reported in BLS' Employment and Earnings Bulletin as of February 1984), were excluded.

Step 4—Within each urban or rural area, the total gross hospital salaries as computed in step 2 were summed for all hospitals not excluded in step 3 to yield the total gross hospital salaries in each area.

Step 5—The total gross hospital salary result computed in step 4 was divided by the corresponding total number of paid hours in the area to yield an average hourly wage for each urban and rural area.

Step 6—The total inflated gross hospital salaries computed in step 2 for all wages not eliminated due to aberrant wage data were divided by the reported number of total paid hours in these hospitals to obtain the national average hourly hospital wage based on gross salaries. This national average is \$8.52.

Step 7—For each urban or rural area, the hospital wage index value was calculated by dividing the average hourly wage computed in step 5 by the national average hourly wage.

Procedure II: Computation of the 1984

wage index.

Step 1—Each of the non-Federal acute care hospitals subject to the prospective payment system for which 1984 data have been received (including hospitals in Puerto Rico) was classified into its appropriate urban or rural area based on the current urban area definitions used in the prospective payment system.

Step 2-For each hospital, the total gross hospital salaries as reported for hospital fiscal years that began in FY 1984 were inflated from the end of the hospital's cost reporting year through August 31, 1985 using the percentage change in average hourly earnings of hospital industry workers (Standard Industrial Classification (S.I.C.) 806) in BLS' Employment and Earnings Bulletin. This was done to eliminate any distortion in the data caused by differing hospital cost reporting years. (August 31, 1985 was the latest end date for hospital cost reporting years in the data collection.)

Step 3—For each hospital, the inflated gross hospital salaries computed in step

a

2 were divided by the reported number of total paid hours to yield an average hourly wage. Hospitals with an aberrant average hourly wage, which was defined as an average hourly wage either less than \$3.35 (the minimum wage in 1984) or greater than \$23.61 (2½ times the national average hourly wage as computed from the data collected), were excluded.

Step 4—Within each urban or rural area, the result computed in step 2 was summed for all remaining hospitals to yield the total gross hospital salaries in each area.

Step 5—The total gross hospital salary result computed in step 4 was divided by the corresponding total number of paid hours in the area to yield an average hourly wage for each urban or rural area.

Step 6—The inflated gross hospital salaries computed in step 2 for all hospitals not eliminated due to aberrant wage data were divided by the reported number of total paid hours in these hospitals to obtain the national average hourly hospital wage based on gross salaries. This national average is \$9.76.

Step 7—For each urban or rural area, the hospital wage index value was calculated by dividing the average hourly wage computed in step 5 by the national average hourly wage.

Procedure III: Computation of a wage index for all hospitals except those located in Puerto Rico, based on a blend of the 1982 wage index (computed under Procedure I) and the 1984 wage index (computed under Procedure II).

Step 1—Wage index values for each urban and rural area computed using 1984 data (Procedure II, step 7) were matched to the corresponding urban and rural wage index values computed using 1982 data (Procedure I, step 7). For both indexes, areas were classified as urban or rural using the current definitions.

Step 2—A blended wage index value for each urban and rural area was computed by adding the 1982 and 1984 wage index values and dividing the result by 2.

The results obtained in step 2 constitute the wage index values for each urban and rural area.

For hospitals located in Puerto Rico, the wage index values are not the result of a blend, but are instead based solely on 1984 data. We do not have usable 1982 wage data for Puerto Rico hospitals since these hospitals were not subject to the prospective payment system in 1984 and 1985 when we collected the 1982 wage data from prospective payment hospitals.

We received 25 pieces of correspondence that commented on the

hospital wage index. The comments and our responses are discussed below.

Comment: Several commenters objected to the use of the 1984 wage data because, the commenters alleged, the data contained numerous errors and many hospitals' data were missing, omitted, or deleted from the data base used in calculating the 1984 wage index. In addition, the commenters believe that the data were not sufficiently edited or audited, nor were hospitals given the opportunity to examine and validate the 1984 wage data prior to publication of the proposed rule. The commenters also believe that data that appears to be incorrect should be corrected and revised rather than excluded from the calculation of the wage index.

Response: As we indicated in the proposed rule, we deleted from the data base the 1984 survey data from every hospital whose average hourly wage was below the minimum wage (\$3.35) or 21/2 times above the national average hourly wage. However, in addition to this edit, we also identified any hospital that submitted data that were not audited and whose average hourly wage increased more than a specified percentage over the 1982 average hourly wage, or decreased, or were missing from our data base. The list of hospitals identified using this edit was returned to each fiscal intermediary with instructions to verify the wages and hours reported and to follow up on

missing survey data.
This process identified

This process identified 66 hospitals that had not responded to the survey and 365 hospitals that reported unaudited data with large increases or decreases. As a result of this effort, we collected 38 of the missing surveys and made 129 edit changes prior to publication of the proposed rule. Consequently, we believe that the 1984 wage data base is complete, with 99.5 percent of all hospitals subject to the prospective payment system responding to the survey. In addition, with respect to the accuracy of the data, it should be noted that the survey data from 66 percent of the hospitals reporting were based on audited cost reports, and that a majority of the data that were not originally audited were subsequently returned to the fiscal intermediaries for verification and correction.

To ensure the accuracy of the data further, we identified those areas in which the 1984 wage index value increased or decreased more than eight percent when compared to the 1982 wage index value. In these instances, we compared the salaries reported on the 1984 survey with the same data elements reported on the cost reports as contained in our Hospital Cost Report

Information System (HCRIS). Significant variations were investigated and resolved. As a result of the edits and the receipt of additional surveys since the proposed rule was published, 22 area wage index values (9 rural and 13 urban) have been revised.

We believe that we have taken all the steps possible to ensure the accuracy of the 1984 wage data. We further believe it is appropriate to exclude data that could not be corrected and that fell outside our minimum and maximum parameters. If we had continued to use these data, the wage index values could have been inappropriately skewed.

As indicated above, certain commenters believe that we should have afforded each hospital the opportunity to examine and verify its data, as we did after we had collected the 1982 data. Since the 1982 data collection was the first we had undertaken, we believed it appropriate to afford hospitals the opportunity to examine and verify the data, particularly since many hospitals were probably not aware of how the data were to be used, and of the necessity for accuracy. These same circumstances do not exist with respect to collection of the 1984 data. Hospitals are or should be aware that we periodically collect data for purposes of developing a wage index. A training session was also held with the fiscal intermediaries, at which the wage data collection effort was discussed, and its importance stressed.

Comment: Four commenters expressed concern that the data reported in the 1984 wage data should include items such as home office salaries, salaries of contract personnel, and salaries of personnel working in related organizations, as defined in Medicare regulations. These commenters argued that the inclusion of these items would result in index values that more closely approximate relative wage differences among areas.

Response: The 1984 survey data represent an update of the 1982 wage data. Wage index values constructed from these data use wages and salaries reported by hospitals in the total wages and salaries column of the trial balance on the Medicare cost report. In addition, we used the same definition for paid hours in developing the two wage indexes.

We did not make adjustments to the wage data for contract labor, home office salaries, and related organizations for three reasons. First, many hospitals indicated problems in determining hours associated with the services described above. Second, wages paid to home offices or related organizations do not

necessarily reflect local economic circumstances and wage patterns. Finally, the fees paid to outside organizations (such as contract nursing labor) often reflect overhead as well as salaries and other items.

However, we also recognize that there may be additional refinements that could be adopted in constructing the hospital wage index. We will continue to investigate which refinements and changes might be appropriate.

Comment: Two commenters wanted to know why we did not use the data reported on the HCFA Form 339, Exhibit 7, to develop a revised wage index based on an occupational mix of

employers.

Response: Exhibit 7 of the HCFA Form 339 (Provider Reimbursement Questionnaire) was developed to allow us to assess the feasibility of developing a wage index that takes hospital occupational mix into account. This form was approved for use in connection with cost reports submitted during calendar year 1986. The collection of these data is incomplete, and they still must be analyzed and reviewed in order to determine their suitability for use.

Comment: Two commenters noted changes in the geographic classifications of several hospitals from what was reported using the 1982 wage data. The commenters alleged that the proposed rule did not contain sufficient information to allow them to ensure that

these changes were proper.

Response: In an effort to ensure the accuracy of the location of a hospital, we compared the county location reported on the 1984 wage survey with other data sources including the 1982 wage data file. Any differences or discrepancies in location were resolved by using the hospital's address.

Comment: One commenter suggested that we eliminate the 1984 wage data for any hospital that has subsequently terminated its participation in the Medicare program or ceased operations

altogether.

Response: We do not believe that it is appropriate to eliminate these hospitals from the survey data. To the extent that data are available for such a hospital for the period covered by the survey, it is appropriate for the data to be used, provided that the data pass edit screens.

The basis of the commenters' arguments is that a hospital that ceases operation or terminates its Medicare participation will not be subject to the new wage index. While this is true, the wage index measures relative wage difference from area to area as of a particular time. If a hospital was in operation at the time of the survey, it

was paying wages and salaries. To eliminate its data could conceivably skew the average hourly wages in that area, either to the detriment or the benefit of the other hospitals in the area.

Comment: We received several comments requesting that the wage index be updated on a regular basis.

Response: In principle, we agree with these recommendations that the hospital wage index be updated on a regular basis. However, while we recognize the need for future updates, changes from year to year may not be significant enough to subject hospitals to the additional reporting requirements and paperwork necessary to accomplish a survey and subsequent follow-up on an annual basis. At present, we do not have a process in place for obtaining wage data on a regular basis. However, we will be investigating the necessity and feasibility of such a process for future updates.

Comment: Two commenters stated that the wage survey data for Puerto Rico are questionable. Specifically, the commenters are concerned that the rural wage index values are higher than most of the urban wage index values, which is contrary to the trend in the 50 States. Additionally, these commenters were concerned as to whether hospitals in Puerto Rico were classified properly as

urban or rural.

Response: We can only speculate as to why the rural hospitals in Puerto Rico experience higher wage levels than the hospitals located in urban areas. The wage index values published in the proposed rule for Puerto Rico based on 1984 data were calculated in the same manner as the 1984 wage index values for the rest of the United States. The wage values for Puerto Rico were derived from the survey data received from the fiscal intermediary for all Puerto Rico hospitals. The wage index values merely reflect the results of the survey.

Nonetheless, as a result of comments received, we discovered that several Puerto Rico hospitals were incorrectly classified as rural hospitals. As a result of making the corrections, the wage index values for several Puerto Rico areas have changed. In addition, as a result of reclassifying hospitals into the correct areas, the Puerto Rico standardized amounts have also changed. Currently, there are only eight Puerto Rico hospitals that are classified as rural, therefore only these few hospitals were used to calculate the Puerto Rico rural standardized rates. The proposed Puerto Rico rates reflected classification of 15 hospitals as rural.

Comment: One commenter wanted to know what inflation factors were used to inflate the wage and salary data used in computing the wage index to August 31, 1985 (see Procedure II, Step 2, above). In addition, the commenter objected to the use of one inflation factor. The commenter indicated that different areas and different hospitals may experience difference rates of wage and salary increases.

Response: The rates used to inflate the wages and salaries reported in the 1984 wage index survey were 5.3 percent for 1984, and 5.3 percent for 1985. These rates were based on the Annual Rate of Change for Average Hourly Earnings for hospital workers (S.I.C. 806) published by the Bureau of Labor Statistics.

With respect to our use of a single inflation factor to bring all the wage data to a common point in time, we had serveral options at the time we begun using the 1984 wage data to create a wage index. One option was not to apply any inflation factor at all. This could have resulted in an index that did not represent relative differences in wages between areas due to the fact that wages for many hospitals could have been low simply because the data were from an earlier period. On the other hand, to reflect the impact of wage inflation among hospitals most accurately, we would have had to require that all hospitals report their gross wages and salaries, along with paid hours, on a monthly basis so that we would be able to track individual hospital inflation rates. Such a system, however, would be administratively burdensome for both HCFA and the hospital industry. Similarly, it would also be burdensome to require all hospitals to report the data for a standard time period, without regard to hospital cost reporting periods.

Not only would these alternative methods have been burdensome, but it has not been demonstrated, beyond the assertion of the commenter, that using a single national inflation factor would in fact lead to significant inaccuracies in the wage data and in the index values. We believe that our use of a single inflation factor represents a reasonable course of action given the alternatives discussed above.

Comment: We received comments on blending the 1982 and 1984 HCFA wage indexes from several commenters. Some commenters agreed that we should implement a blended rate to lessen the impact of large changes in the index values for some areas. However, one commenter did not believe that such a practice would be appropriate for more then one year. Other commenters recommended that the current wage index be retained until a more current

survey could be validated by hospitals and a means of annually updating the wage index established. All commenters found the 1982 data preferable to the 1984 data for various reasons although there were concerns about the age of the 1982 data. While recognizing that blending would smooth out the fluctuations that would result from moving to an index based on 1984 data, the commenters would prefer that we continue to use the 1982 data until better and more recent data are available.

Response: We have responded to the criticism of the accuracy of the 1984 HCFA wage data above and believe that we have taken all reasonable measures to ensure the accuracy of the data. Since we have established what we believe to be a reliable wage index, based upon the most recent data available, it is appropriate to begin using the new wage index. However, we do not believe that it would be prudent on our part to implement this change without providing some mechanism for protecting hospitals from abrupt changes in their payment amounts. To continue to use the 1982 wage index until further validation of the 1984 data can be accomplished, and a procedure developed for annually updating the wage index, would not be appropriate. since we would not be using the most recent data available for purposes of computing the prospective payment

The principle of using the most recent data available was firmly established by Congress at the time the prospective payment system was first instituted. In addition, administrative law and judicial rulings do not require that changes or improvement to a system be perfect before they are adopted; rather, the criteria that should guide an agency's deliberations is one of 'reasonableness," that is, has the agency acted reasonably or taken a reasonable approach in adopting such changes or improvements. We believe that we have met this standard both in the development of the 1984 wage index and the adoption of a blended wage index for payment purposes.

IV. Inclusion of Puerto Rican Hospitals in the Prospective Payment System

When section 1886(d) was added to the Act by Pub. L. 98–21, all hospitals located outside the 50 States and the District of Columbia were excluded from the prospective payment system and thus have continued to be paid on the basis of reasonable costs subject to the rate-of-increase limits established by section 1886(b) of the Act. However, section 9304(a) of Pub. L. 99–509 added a new section 1886(d)(9) to the Act to

include eligible Puerto Rico hospitals in the prospective payment system effective with discharges occurring on or after October 1, 1987.

Section 1886(d)(9)(A) following (ii) of the Act specifies that a hospital is subject to the prospective payment system if it is located in Puerto Rico and otherwise would be subject to that system if it were located in one of the 50 States. Although eligible Puerto Rico hospitals are to be included in the prospective payment system, there are some special rules that apply to those hospitals.

Section 1886(b)(9)(A) of the Act specifies that the payment per discharge under the prospective payment system for hospitals in Puerto Rico is the sum of—

- 75 percent of the Puerto Rico discharge-weighted urban or rural standardized rate.
- 25 percent of a national dischargeweighted standardized rate.

We proposed to compute separate urban and rural standardized payment rates for Puerto Rico. For FY 1988, section 1886(d)(9)(B)(i) of the Act specifies that this computation is to be done in the same manner we used to compute the regional standardized rates under section 1886(d)(2) of the Act, except that the rate is to be based on the Puerto Rico hospitals' target amounts (as defined in section 1886(b)(3)(A) of the Act) that were applicable for cost reporting periods beginning on or after October 1, 1986, updated to the midpoint of FY 1988 by prorating the applicable percentage increase (that is, the percentage increase in the market basket index minus 2.0 percentage points). Under section 1886(d)(9)(A)(ii) of the Act, the national standardized rate that makes up 25 percent of the payment rate for Puerto Rico hospitals consists of the discharge-weighted average of the national rural standardized amounts and the national urban standardized amounts that are used for paying all prospective payment hospitals outside of Puerto Rico.

As required by section
1886(d)[9)[B)(vi) of the Act, the laborrelated portion of the Puerto Rico
standardized amount is adjusted by the
appropriate wage index value for the
area in which a Puerto Rico hospital is
located. We proposed to include Puerto
Rico in the HCFA wage index that is
used for all prospective payment
hospitals and to adjust the Puerto Rico
standardized amount for each area to
reflect the average wage level relative to
the national average wage.

For FY 1989 and subsequent fiscal years, section 1886(d)(9)(C)(i) of the Act

specifies that the Puerto Rico standardized amount is to be updated by the applicable percentage increase determined by the Secretary under section 1886(e)(4) of the Act. Section 1886(e)(4) of the Act further specifies that the update factor applied to Puerto Rico hospitals must be the same as the update factor applied to prospective payment hospitals located in the 50 States and the District of Columbia.

Section 1886(d)(9)(D) of the Act specifies that the following provisions of section 1886(d)(5) of the Act concerning additional payments to, or special treatment of, prospective payment hospitals also apply to prospective payment hospitals in Puerto Rico:

 Section 1886(d)(5)(A) of the Act, which requires that additional amounts be paid for outlier cases.

 Section 1886(d)(5)(B) of the Act, which requires that additional amounts be paid for indirect medical education costs.

• Section 1886(d)(5)(c)(iii) of the Act, which authorizes the Secretary to make other exceptions and adjustments as the Secretary deems appropriate.

 Section 1886(d)(5)(E) of the Act, which permits payment on a reasonable cost basis for anesthesia services furnished in a hospital by a certified registered nurse anesthetist (CRNA).

 Section 1886(d)(5)(F) of the Act, which authorizes additional payment for hospitals that serve a disproportionate share of low-income patients.

The following provisions of section 1886(d)(5) of the Act do not apply to prospective payment hospitals in Puerto Rico:

- Special treatment of referral centers (section 1886(d)(5)(C)(i) of the Act).
- Special treatment of sole community hospitals (section 1886(d)(5)(C)(ii) of the Act).

We proposed that the following types of hospitals and hospital costs that receive special treatment in the prospective payment system under section 1886(d)(5)(C)(iii) of the Act would also receive special treatment if located in Puerto Rico:

- Hospitals involved extensively in treatment for and research on cancer that meet the requirements of § 412.94.
 - Christian Science Sanatoria.
- Hospitals that are located in urban areas and that are reclassified as rural, as described in § 412.104.
- Hospitals with a high percentage of discharges for end-stage renal disease patients, as described in § 412.104.
- Hospitals approved as renal transplantation centers.
- Hospitals in redesignated rural counties that are surrounded on 95

percent of their perimeters by urban counties, as described in § 412.63(b)(3).

We proposed to add a new Subpart K to Part 412 to implement the special rules that apply to prospective payment hospitals located in Puerto Rico.

Conforming changes were also made in §§ 412.23(f).

Section 1886(e)(1)(C) of the Act, as added by section 9304(c) of Pub. L. 99–509, requires that for discharges occurring in FY 1988, the aggregate payment to prospective payment hospitals including those hospitals located in Puerto Rico be equal to the aggregate payment that would have been made to those hospitals under prior law; that is, the addition of hospitals in Puerto Rico to the prospective payment system must be "budget neutral".

As explanation of the methodology used to calculate the payment rates for hospitals in Puerto Rico, as well as the budget neutrality issue, is set forth in sections III and IV of the addendum to this final rule.

Comment: One commenter wanted to know how the labor-related and nonlabor-related components of the Puerto Rico standardized amounts were determined. Another commenter questioned whether the same labor and nonlabor portions were used in determining both the Puerto Rico standardized amounts and the national standardized amounts.

Response: As indicated in the proposed rule (52 FR 22107), the labor and nonlabor portions of the target amounts were based on the labor and nonlabor components of the hospital market basket. The latest hospital market basket components were published in the September 3, 1986 final rule (51 FR 31530). Based on these market basket components, the labor portion represents 74.39 percent of the Puerto Rico target amounts and the nonlabor portion is 25.61 percent. These are the same portions that are used in determining the national standardized amounts.

The source of the second commenter's confusion concerning the labor and nonlabor portions reflects a misunderstanding as to how the labor and nonlabor portions are determined for purposes of computing the standardized amounts. The commenter believes that a national average cost per discharge (or, in Puerto Rico, target amount per discharge) is first determined, and that this overall average is then split into labor and nonlabor portions. However, in actuality, each individual hospital's cost per discharge (for hospitals outside Puerto Rico) or target amount per

discharge (in the case of Puerto Rico) is divided into labor and nonlabor portions using the constant percentages discussed above. Then, separate national and regional averages are computed for the labor portion, after the standardization for different area wage levels, and the nonlabor portion respectively.

respectively.

Comment: One commenter requested that we publish the case-mix index values for Puerto Rico hospitals that were used to standardize the Puerto Rico prospective payment rates. The commenter noted that the Puerto Rico hospital case-mix index values were not included in table 3c of the June 10, 1987 proposed notice along with the case-mix index values of all other prospective payment hospitals.

Response: We note that the case-mix index table published in the proposed notice represents FY 1986 case-mix index values and was published for the sole purpose of identifying hospitals that may qualify as rural referral centers. Since the rural referral center provisions does not apply to Puerto Rico hospitals, their case-mix index values were not

published.

As indicated in the proposed rule (52 FR 22107), the case-mix index values used to standardize the Puerto Rico rates were from FY 1984. These data, as well as all other data used to compute the standardized amounts, are available to the public upon request. Generally, we have not published all data used to compute prospective payment rates because of the volume and magnitude of the data.

Comment: One commenter stated that the same cost outlier thresholds applicable to prospective payment hospitals located outside of Puerto Rico are not appropriate for Puerto Rico hospitals given the fact that hospital costs are lower in Puerto Rico. This commenter suggested that cost outlier thresholds for Puerto Rico should be based on actual Puerto Rico hospital cost experience. However, another commenter stated that Puerto Rico hospitals have higher costs for some items than do other hospitals and that, therefore, they should receive a cost-ofliving adjustment similar to the adjustment for hospitals in Alaska and

Response: Section 1886(d)(9)(D) of the Act (as added by section 9304 of Pub. L. 99–509) specifies that certain provisions (including outlier payments but not a cost-of-living adjustment) applicable to subsection (d) hospitals "shall apply to subsection (d) Puerto Rico hospitals * * * in the same manner and to the extent as they apply to subsection (d) hospitals * * *." Therefore, we are

using the same day and cost outlier thresholds for Puerto Rico hospitals and all other hospitals. However, when determining whether a case qualifies as a cost outlier, the threshold amount is adjusted by the hospital's wage index value. In effect, this results in the outlier formula taking into account the specific cost experience of the hospital.

We note that, under current outlier policy, which we will not be revising as a part of this final rule (see section V.B. of this preamble), the preponderance of outlier cases will be paid as day outliers, thus reducing the impact of using uniform cost outlier thresholds. Further, we note that there are also differences in hospital costs among different regions located outside of Puerto Rico, but that different outlier thresholds were never provided in recognition of these differences.

Finally, we note the commenter's justification in support of the commenter's position that Congress recognized Puerto Rico's special cost situation by providing unique standardized rates for Puerto Rico. However, Congress also incorporated features that are identical to the features applicable to the prospective payment system for all hospitals outside of Puerto Rico. In fact, Congress provided that Puerto Rico hospitals will be entitled to additional payments for the indirect costs of medical education and as disproportionate share hospitals, even though the formulas for computing these adjustments would be different (and perhaps result in lower adjustments) if they were based solely on Puerto Rico data and circumstances.

For these reasons, and in view of the disagreement among the commenters as to whether Puerto Rico hospitals have higher or lower costs, we are not adopting at this time the suggestion that we develop outlier thresholds specific to Puerto Rico.

Comment: One commenter expressed concern that we have not appropriately identified all hospitals in Puerto Rico that qualify for a disproportionate share adjustment. The commenter also indicated that several hospitals in Puerto Rico would qualify for a disproportionate share adjustment under § 412.106(b)(2) because they receive at least 30 percent of their inpatient revenue from the Commonwealth government for the care of indigent patients.

Response: We should point out that the data we used to standardize the Puerto Rico payment amounts to reflect the disproportionate share adjustment were based on FY 1984 cost report data as well as Medicaid data supplied by the Department of Health in Puerto Rico. These data were used solely for the purpose of adjusting the Puerto Rico rates. The determination of whether a hospital is entitled to receive additional payments as a disproportionate share hospital is made by the fiscal intermediary based on the latest data available.

The determination of whether a hospital would qualify for a disproportionate share adjustment under § 412.106(b)(2) would be made following the end of the hospital's cost reporting period, at which time the hospital will have the opportunity to apply for an adjustment under the provision on revenue from State and local governments for indigent care. Since we have no data concerning which hospitals would qualify for a disproportionate share adjustment under this provision, no adjustments were made to the Puerto Rico rates to reflect payments under this provision. However, any hospital that can demonstrate to its intermediary that it does, in fact, qualify for this adjustment, will receive disproportionate share adjustments as appropriate.

Comment: One commenter requested that hospitals in Puerto Rico be allowed to appeal their rate-of-increase target amounts used to compute the Puerto Rico standardized amounts. The commenter contends that prospective payment hospitals outside of Puerto Rico were permitted to appeal their base period costs upon entering the prospective payment system and, therefore, hospitals in Puerto Rico should be afforded the same opportunity to challenge their target amounts.

Response: The circumstances under which hospitals in the fifty States and the District of Columbia were afforded the opportunity to provide additional cost report data involved the computation of the hospital-specific portion of the prospective payment rates during the transition period to a fully Federal rate. However, since there is no transition period provided under the law for Puerto Rico hospitals under the prospective payment system, hospitals in Puerto Rico will not have a hospitalspecific portion in their prospective payment rate, but rather will be paid using fully Federal standardized payment amounts (that is, 75 percent Puerto Rico Federal rate and 25 percent national Federal rate). We should point out that at the time the target amounts were computed, hospitals in Puerto Rico also had the opportunity to appeal their base period cost report data used to compute the target amounts.

The Federal standardized payment amounts for prospective payment

hospitals outside of Puerto Rico were computed using 1981 unaudited cost report data. These data represented the latest cost data available to us at the time the initial standardized rates were computed. Hospitals were not given the opportunity to submit additional data for these 1981 cost reports. Likewise, the rate-of-increase target amounts we used to compute the Puerto Rico standardized amounts were the latest available to us at the time. While target rates for particular years may be revised under existing regulations for purposes of determining the amounts of payment for those years under the reasonable cost reimbursement principles, to permit their revision to affect the Puerto Rico standardized amounts would, in effect, defeat Congress' intention that the standardized amounts be prospectively determined based on the best data available.

As we stated in the proposed rule, if we were to allow constant revision of the Puerto Rico standardized amounts based on changes to the hospitals' target rates, we would create continuing uncertainty as to what the prospective rates are. Also, for years after FY 1988, section 1886(d)[9](C)(i) of the Act requires that the previous year's Puerto Rico standardized amounts be updated by the applicable percentage increase determined for the prospective payment system. We do not believe that Congress contemplated changes in those amounts because of revisions in the data base.

Furthermore, revising the Puerto Rico standardized amounts to take into account revisions in target rates would be contrary to our policy that we not make changes to the standardized amounts because of changes to the data base used to calculate the standardized amounts. We believe our policy is in accordance with congressional intent to use the best data available. We note that we did not revise the original prospective payment standardized amounts that were effective October 1. 1983 to take into account revisions in the data that were used to calculate those amounts (that is, cost reports for reporting periods ending in calendar year 1981). Therefore, we would allow revisions in the target rates for individual cost reporting periods subject to the rate-of-increase limits under the current regulations for purposes of determining payment for those periods. However, these revisions would have no impact on the data used for the computation of the Puerto Rico standardized amounts.

V. Other Decisions and Changes to the Regulations

A. Review of DRG Assignments (§§ 412.60 and 466.70)

We have encountered situations in which a hospital that submits a claim to Medicare for payment later attempts to resubmit the claim based on additional information that would place the case in a higher-weighted DRG. Some corrections of billing information are warranted if, for example, the hospital omitted critical documentation or misread the medical record. We believe that it is appropriate to allow a hospital a reasonable period of time in which to correct its own error by submitting additional or corrected information on an adjustment bill. Nevertheless, as in the case of any business transaction, we do not believe it is appropriate for the billing party to revise a claim long after the original claim is submitted and paid.

Allowing hospitals an extended period of time to discover errors and to resubmit bills is contrary to good business practice. A workable prospective payment system would not exist if the fiscal intermediaries are constantly processing recoded claims based upon the same documentation or if bills lack finality because they are forever subject to revision.

Therefore, effective April 23, 1964, we established an informal review mechanism through administrative directive by issuing changes to the following manuals:

- Hospital Manual (HCFA Pub. 10), section 287.5, transmittal number 382.
- Medicare Intermediary Manual (HCFA Pub. 13–3), section 3798, transmittal number 1109.

These issuances specified that a hospital has 60 days after the date of an initial DRG assignment to a claim to request review. The hospital may submit additional information as a part of its request. The fiscal intermediary reviews the data and adjusts the DRG if appropriate.

As part of the PRO's review responsibility, the initial PRO contract cycle provided for review of hospital requests for DRG claims adjustments submitted after the initial claim had been filed. This review applied only if the intermediary's review resulted in the assignment of a higher-weighted DRG and the PRO had not previously reviewed the case in question. Because these claims adjustments were considered to represent a high risk of DRG manipulation, 100 percent of these cases were reviewed postpayment. The PRO not only determines if the request for coding changes is appropriate, but

also conducts full PRO review of the case if this review was not performed previously. The PROs collected data on the frequency with which hospitals submitted erroneous requests for DRG claim adjustments. Identification by the PRO of a pattern of inappropriate coding adjustments required corrective actions. The second PRO contract cycle effective July 1, 1986, required that this review be conducted on a prepayment basis.

We proposed to include the provisions of the manual instructions concerning hospitals' requests for review of DRG assignments in the regulations. We proposed to review § 412.60 to specify that a hospital has 60 days to request a review by the intermediary of a DRG assignment and to describe how that review is conducted. In addition, we proposed to revise § 466.70 to provide that a PRO must review every case in which a higher-weighted DRG is assigned to a discharge as a result of the

intermediary's review.

Comment: A number of commenters were not in favor of the proposal to codify the 60-day limitation on submittal of requests for review of the DRG initially assigned to a claim by the fiscal intermediary. More specifically, most commenters opposed the imposition of any limitation on the time allowed for a hospital to submit revised or corrected coding information to the fiscal intermediary for purposes of obtaining a more accurate DRG assignment. They believe that it is unnecessary to set a time limit because hospitals already have clear incentives to request promptly correction of DRG assignments that lead to inappropriately low payments; and conversely, the Medicare program is not harmed but actually benefits financially from any delay in hospitals' requesting corrections. Their complaint was that a time limit would only penalize those hospitals that, because of circumstances beyond their reasonable control, cannot always identify and prepare corrections within

The commenters were concerned that the 60-day limitation will prevent hospitals from conducting their own review in order to identify incorrect or inadequate coding. Many commenters believe that hospitals should have the right to correct any coding mistake discovered on any bill processed by the fiscal intermediary in the last 90 to 180

days.

Response: We believe that the fact that the majority of the commenters are apparently unaware of the fact that the 60-day limitation on submittal of revised or corrected codings has been our stated policy since April 23, 1984, justifies our decision to include both the review by

the fiscal intermediary and the review by the PRO in the regulations. Initially, requests for corrections of coding were processed as adjustment bills under procedures that were established prior to the implementation of the prospective payment system. As more hospitals became subject to that system, the hospitals became aware of the importance of correct coding. Hospitals came to appreciate that some cases could have been coded more advantageously under allowable International Classification of Diseases-9th Edition-Clinical Modification (ICD-9-CM) protocols. (The coding system of the ICD-9-CM is the one on which DRG assignments are based.)

The 60-day limitation was included in manual instructions to provide a reasonable control on the volume of corrections that would have to be processed. Since the manual issuances specified that hospitals could present additional information to the fiscal intermediary in the form of corrected coding, we expected that requests for review of DRG assignments might largely consist of corrections that would result in the assignment of a higherweighted DRG (that is, that it might result in "upcoding"). However, corrections resulting in higher-weighted DRGs were made subject to PRO review (if not previously reviewed) to guarantee that the new coding was appropriate in view of the medical record of the case. .

We agree that hospitals should have a reasonable amount of time to correct errors. However, we are concerned about a review process that, while ostensibly having as its goal only complete and accurate coding, generally results in greater numbers of higherweighted DRGs as opposed to equal numbers of higher and lower-weighted DRGs. Setting a time limit on revision of a claim serves the dual purpose of providing hospitals a reasonable period of time to correct errors and of ensuring that claims are considered final for payment purposes on an ongoing basis.

Comment: Commenters believe that PRO review of those cases resulting in assignment in a higher-weighted DRG provides adequate controls to ensure that payment of DRGs is not manipulated due to rebilled claims and that therefore a 60-day limit on requesting corrections is not necessary.

Response: Even assuming that the PRO review were adequate to prevent manipulation, as noted above, the 60day limit also serves to provide a reasonable control on the volume of corrections that would have to be processed.

Comment: One commenter agreed that allowing hospitals or HCFA an extended period of time to discover errors and resubmit bills contradicts good business practice. However, the commenter suggested that a 90-day limit would be more realistic and would permit hospitals to include corrected bills in their cost reports, which are due 90 days after the close of the cost reporting period. Another commenter believes that it is inappropriate to compare Medicare claims with common business practice because the prospective payment system does not base payments on fee-for-service, as do most businesses, but on a complicated coding system. The commenter suggested that a period of 180 days to submit changes would allow prudent hospitals time to use their audit and review procedures to discover errors. The need for this time was linked to the high potential for errors in coding due to ongoing changes in the ICD-9-CM and high turnover in medical records personnel. The need to obtain physician attestation on bills was also cited.

Response: We appreciate the support of the first commenter in our goal of maintaining a current, reliable billing process. We believe that a 60-day limitation, beginning with the date of the fiscal intermediary's payment notification, is a sufficient amount of time in which to uncover errors in coding. Although it is true that the ICD-9-CM coding process is complicated and subject to ongoing changes, we believe that the combined efforts of the American Medical Records Association, the American Hospital Association. National Center for Health Statistics, and HCFA have provided an invaluable resource for medical records personnel in resolving coding questions.

Since the physician attestation must be obtained prior to submission of the bill to the fiscal intermediary, the need to obtain attestations should not affect resubmission of claims involving coding errors. Revised attestations are only needed when diagnoses or procedures in the original attestation must be changed.

Comment: Several commenters argued that the 60-day limitation was especially unfair because the fiscal intermediaries and PROs have no time limitations on the adjustment or review of claims. Commenters also recommended that PROs and fiscal intermediaries should be given more specific instructions concerning the review of claims, and that hospitals should have a method of recourse if the request for a change in DRG is denied.

Response: We believe that the commenters' concerns could have been partially allayed with a more complete description in the proposed rule of the existing instructions to the fiscal intermediary and PROs. We assumed that this was an area already widely understood by a substantial number of hospitals, and we cited only the references in the manual that impose the 60-day limitation rather than describing the procedure in detail. (See 52 FR 22089.) The instructions for fiscal intermediaries on processing adjustment bills can be found in section 3816.1 of the Part A Intermediary Manual (HCFA Pub. 13-3). Prior to implementation of the prospective payment system, and for some time thereafter, the number of hospital-initiated adjustment bills under the provisions of section 3816.1 was very low, accounting for a small percentage of the fiscal intermediaries' processing time. By March 1984, a number of hospitals were submitting adjustment bills with corrected coding, creating an unaccustomed workload. Our estimate of the potential increase in this workload led us to impose a 60-day limitation on those claims submitted for DRG adjustment. Since claims had already been paid, these bills and subsequent submittals under the 60-day rule have been given a relatively low priority for processing.

We appreciate the comments pointing out the potential processing lag at the fiscal intermediary level. We will continue to measure the actual impact of these adjustment bills and will evaluate the need to monitor fiscal intermediary

performance in this area.

PROs, on the other hand, have specific instructions and processing deadlines both in their contracts with HCFA and in the instructions outlining this review and they are reviewed by the regional office to ensure compliance with these requirements. (See the PRO Manual (HCFA Pub. 19), sections 2003, 2004, and 2050.4E.) All medical review activities included in section 2050 of the PRO manual must be instituted (that is, cases identified and records requested) within 15 calendar days of the receipt of the intermediary data.

The review of the case must be completed within 15 calendar days of the receipt of the medical record. Under section 2050.4E of Pub. 19, in order to complete the review of a revised DRG, the PRO must be supplied with the following information from the hospital:

- · The initial codes submitted.
- The codes submitted for adjustments.
- A statement explaining why the original codes were submitted incorrectly.
 - · A copy of the medical record.

 If coding changes were based on newly acquired clinical information, a copy of such information (for example,

an autopsy report).

We will continue to monitor the processing of hospital requests for DRG adjustment to determine whether the omissions and errors made by coding personnel justifies additional training efforts by the ICD-9-CM Coordination and Maintenance Committee. Although the processing of these bills by fiscal intermediaries will continue to have a lower priority than initial payment processing and coordination, we will monitor the number of these requests to determine if there is a substantial adverse impact of the procedure on the overall Medicare payment to any one hospital or any group of hospitals and make any necessary adjustments.

A hospital that receives an unfavorable decision on its request for review of DRG assignment may appeal that decision under the provisions of 42

CFR Part 405, Subpart R.

B. Increase in the Prospective Payment Rates and Rate-of-Increase Limits (§§ 412.63, 412.73, and 413.40)

Section 9302(a)(1) of Pub. L. 99-509 amended section 1886(b)(3)(B)(i)(II) of the Act to provide that the applicable percentage increase for FY 1987 is 1.15 percent and for FY 1988 is the market basket percentage increase minus 2.0 percentage points. (We note that the Congress is considering modifying the applicable percentage increase for FY 1988, so that the market basket change minus 2.0 percentage points may not be the final update factor.) A final rule published in the Federal Register (52 FR 42229) on November 24, 1986 amended §§ 412.63, 412.73, and 413.40 to implement the changes applicable to FY 1987. We proposed to amend those same sections to implement the provisions of section 1886(b)(3)(B)(i)(II) of the Act applicable to FY 1988. We received no comments on this proposal.

C. Payment for Outlier Cases (§§ 412.82 and 412.84)

Section 1886(d)(5)(A) of the Act requires that, in addition to the basic prospective payment rates, payments must be made to hospitals for atypical cases known as "outliers". These are cases that have either an extremely long length of stay or extraordinarily high costs when compared to the other discharges classified in the same DRG.

Section 1886(d)(5)(A)(iii) of the Act specifies that the outlier payments should approximate the marginal cost of care beyond the outlier threshold. In the September 1, 1983 interim final rule, we established the ratio of marginal cost to

average cost at 60 percent (48 FR 39776). Therefore, the regulations (§§ 412.82 and 412.84) currently provide that the marginal cost of outlier cases is based on a 60 percent factor.

For day outliers, an additional per diem payment is made for each covered day of care beyond the threshold. The per diem payment is based on 60 percent of the average per diem Federal rate for the DRG, which is calculated by dividing the wage-adjusted Federal rate for the DRG by the geometric mean length of stay for that DRG. During the transition period (cost reporting periods beginning on or after October 1, 1983 and before October 1, 1987), this amount is multiplied by the applicable Federal blend percentage. Starting with cost reporting periods beginning on or after October 1, 1987, the Federal portion is 100 percent of the payment rate.

For cost outliers, the additional payment is based on 60 percent of the difference between the hospital's adjusted charges for the discharge and the outlier threshold. We determine the cost of the discharge based on 66 percent of the billed charges for covered services. The cost is further adjusted to exclude an estimate of indirect medical education costs and payments for hospitals that serve a disproportionate share of low-income patients. As with day outliers, the resulting amount is then multiplied by the applicable Federal

portion of the blend.

Our analysis indicates that while our payment policy for outliers effectively reduces the risk faced by hospitals in treating cases that are outside the normal range of cases in terms of care or costliness, additional compensation would be justified for the most expensive case, particularly those long-stay cases with extremely high costs. On the other hand, some cases that qualify for additional payment as day outliers are not extraordinarily costly. Therefore, we proposed to make two changes to the

outlier regulations.

We currently pay even the most expensive day outliers at a per diem amount that is based on the average payment for all discharges assigned to that DRG. For some of the cases that currently qualify as day outliers (and therefore cannot be cost outliers), the per diem rate paid for those cases does not adequately compensate the hospital for its costs. This is especially true in day outlier cases with extremely high costs (for example, severe burn cases) for which the daily costs vastly exceed the day outlier per diem and for which that daily difference is multiplied by a long length of stay. Thus, we proposed that if a day outlier case also meets the

cost outlier criteria, we would pay the case using the cost outlier methodology. Day outliers that do not meet the cost outlier criteria would continue to be paid on the basis of a per diem rate.

We also reexamined the 60 percent marginal cost factor used in calculating the payment for outlier cases. Evidence from recent research indicates that a higher marginal cost factor would result in more appropriate payments for the most expensive cases by more effectively reducing the financial risk to hospitals that is associated with these cases. In particular, we note that the estimated loss per case is higher for cost outliers than for day outliers. When day outliers are separated into two categories-those exceeding the day outlier threshold but not the cost outlier, and those exceeding both the day and the cost outlier thresholds-the estimated loss per case for the more expensive day outliers (those also exceeding the cost threshold) substantially exceeds that for the less expensive day outliers (those that do not exceed the cost threshold).

This finding that the financial risk associated with outlier cases varies substantially with whether the case is long stay only or exceeds the cost threshold suggests that the marginal cost factor we are using for the most expensive outliers (those that exceed the cost outlier threshold) is too low. Accordingly, for discharges occurring on or after October 1, 1987, we proposed to set payment for the most expensive outlier cases (that is, all outlier cases exceeding the cost outlier threshold) at 80 percent of adjusted charges beyond the cost outlier threshold. Based on our research to date, we believe that this revised marginal factor would result in more adequate compensation to hospitals treating the most costly outlier

cases. Comment: We received a number of comments concerning our proposed changes in the outlier payment policy. While most commenters supported an outlier policy that pays a higher fraction of outlier payments for extremely costly cases, many were concerned about the impact of the proposed changes and recommended that changes in the outlier policy be delayed until further study can be accomplished. The areas of particular concern were the continued use of national cost-to-charge ratios to pay cost outliers as well as the negative impact of the proposed outlier policy on certain groups of hospitals, such as teaching and small rural hospitals.

Response: Given the concerns expressed by commenters, we have decided to delay implementation of any changes to the outlier policy. We are continuing our research concerning the impact of using national ratios in computing cost outlier payments. Preliminary studies conducted since the publication of the proposed rule indicate that, in general, hospitals that have large profits per case under the basic rates have lower cost-to-charge ratios than do hospitals that have smaller profits per case under the basic rates. This means that the use of the national cost-to-charge ratio results in a transfer of payments to hospitals that are doing very well from hospitals that are doing less well.

While the use of hospital-specific cost-to-charge ratios may be more accurate for purposes of computing cost outlier payments, there are a number of significant administrative and data problems associated with using these ratios. For example, estimating future outlier payments in order to establish appropriate outlier thresholds becomes more of a problem, since hospital specific cost-to-charge ratios used in the estimate would not be the same as those used for actually paying outliers. In addition, major changes in PRICER software (the program used to calculate each hospital's payment per discharge) and the Medicare cost report would be necessary in order for cost-to-charge ratios to be developed for payment purposes

Therefore, rather than implement an outlier policy that would place greater emphasis on the cost outlier payment methodology, which could unfairly disadvantage certain hospitals, we have decided to continue with the current outlier policy until we can complete our analysis.

In addition, in response to the commenters' request, and as part of our ongoing analyses of outlier payments, we are continuing to look at the distribution of charges and costs by DRG and type of hospital.

Comment: A few commenters recommended that the outlier pool be increased from five percent to six percent, the maximum allowed under section 1886(d)(5)(A)(iv) of the Act.

Response: Since we did not propose to make any change to the aggregate outlier reduction, we believe it is inappropriate to make such a substantial change at this time without benefit of public comment. Also, since we are continuing, in response to comments, to study which changes might be desirable in outlier payment methodology, rather than implementing our proposed changes for FY 1988, we believe it would be inappropriate to make any changes in the outlier pool until that analysis is completed.

Comments: A few commenters pointed out that outlier payments have consistently fallen short of the outlier reserve and that we have failed to publish data on the amount of outlier payments made since the prospective payment system began. Commenters also stated that we should pay any outlier underpayments from prior prospective payment fiscal years.

Response: We responded to similar comments in the September 3, 1986 final rule (52 FR 31525). In addition, in the Department's report "DRG Refinement: Outliers, Severity of Illness and Intensity of Care," which was submitted to Congress on June 12, 1987, we presented updated outlier payment data for FY 1984 that indicated that actual payments for outliers were about 1.1 percent of total prospective payments, or 53 percent of the 2.1 percent target (5.7 percent of the Federal portion of

prospective payments).

The shortfall in outlier payments for the first year of the prospective payment system was due mainly to the fact that we did not anticipate the steep decline in average length of stay that occurred. As with all other aspects of the system, we used the most recent data available at the time to estimate outlier payments in establishing thresholds. Outlier thresholds in subsequent years were adjusted to reflect the observed decline in length of stay. However, due to a continued decline in length of stay, preliminary outlier payment data for FY 1985 indicate that outlier payments may fall short of the 2.5 percent target for FY 1985 (based on 50 percent Federal portion). We expect the shortfall in FY 1985 to be less than in the first year since thresholds were based on data that included prospective payment experience. As the prospective payment transition period progresses and changes in hospital operations stabilize (as is evidenced by the latest length of stay and case-mix data), we expect that outlier payments will be closer to target.

As required by the law, we estimate, using the most recent data available. what the level of the outlier thresholds must be in order to yield the proper total amount of outlier payments. This is inherently a prospective process and the resulting estimate may be determined to be inaccurate based on later data. However, payment of additional outlier monies based on retrospective adjustments to the thresholds would not be appropriate. Had our original estimates been made to favor hospitals, we would not have later recouped any amounts paid over and above what was set aside in the outlier pool. We note that we do not recoup any part of

payments already made due to subsequent data that indicate upcoding or overstatement of rates.

Comment: Some commenters objected to the fact that in computing the amount of costs for outlier payments, we standardize for indirect medical education and disproportionate share adjustments, which results in reduced outlier payments for teaching and disproportionate share hospitals.

Response: We believe that in computing the amount of the cost outlier payment it is appropriate to standardize costs for the indirect medical education and disproportionate share adjustments. This is because the outlier thresholds represent standardized costs. Since teaching and disproportionate share hospitals generally incur higher costs (and charges) for treating the same cases as nonteaching and nondisproportionate share hospitals, it is appropriate to adjust costs to account for these higher costs before applying the standard outlier threshold. The indirect medical education and disproportionate share adjustments are then appropriately added to the amount of the cost outlier payment.

D. Payments to Sole Community Hospitals (§ 412.92)

Section 1886(d)(5)(C)(ii) of the Act requires that the special needs of sole community hospitals (SCHs) be taken into account under the prospective payment system. The statute specifies a special payment formula for hospitals so classified and further provides for additional payment to SCHs experiencing a significant volume decrease (that is, more than a five percent decrease in total discharges of inpatients) because of circumstances beyond their control. The statute defines SCHs as those hospitals that, by reason of factors such as isolated location, weather conditions, travel conditions, or absence of other hospitals (as determined by the Secretary), are the sole source of inpatient hospital services reasonably available to Medicare beneficiaries in a geographic area. Regulations governing the special treatment of SCHs under the prospective payment system are set forth in § 412.92.

Currently, § 412.92(e) provides that an SCH is eligible for a payment adjustment in any cost reporting period if it experiences more than a five percent decrease in its total discharges for inpatients as compared to its immediately preceeding cost reporting period. To qualify for a payment adjustment, an SCH must submit documentation demonstrating the size of the decrease and the resulting effect on per discharge costs, and show that the

decrease is due to extraordinary circumstances beyond the hospital's control, such as strikes, fires, floods, earthquakes, inability to recruit essential physician staff, or unusually prolonged severe weather conditions.

We determine on a case-by-case basis whether an adjustment will be granted and the amount of that adjustment. As specified in § 412.92(e)(3), a per dicharge payment adjustment, including at least an amount reflecting the reasonable cost of maintaining the hospital's necessary core staff and services, is determined based on the individual hospital's needs and circumstances, the hospital's fixed and semi-fixed costs not paid on a reasonable cost basis, and the length of time the hospital has experienced a decrease in utilization.

Based on our experience with this provision and the applications we have received from SCHs for a volume adjustment, we believe that it is appropriate at this time to clarify the regulations at § 412.92(e). Section 1886(d)(5)(C)(ii) of the Act provides that if an SCH experiences a decrease of more than five percent in its total number of inpatient cases due to circumstances beyond its control, "* * * the Secretary shall provide for such adjustment to the payment amounts under this subsection * * * as may be necessary to fully compensate the hospital for the fixed costs it incurs in the period in providing inpatient hospital services, including the reasonable cost of maintaining necessary core staff and services." We believe that this language makes it clear that a hospital that has continued to receive payments under the prospective payment system that are greater than its inpatient operating costs, even though there has been a decline in occupancy, is not entitled to receive a payment adjustment. Hospitals that receive payments that are greater than the hospitals' Medicare inpatient operating costs have been "fully compensated" for those costs by the prospective payment system. Consequently, we believe that no further adjustment should be granted to these hospitals. Therefore, we proposed to revise § 412.92(e)(3) to make it clear that any adjustment amounts granted to SCHs for a volume decrease may not exceed the difference between the hospital's Medicare inpatient operating costs and total payments made under the prospective payment system, including outlier payments and indirect medical education payments.

We also proposed to revise § 412.92(e)(2)(ii), which currently requires that, in order to receive a volume adjustment, the decline in the hospital's total discharges must be due to extraordinary circumstances beyond the hospital's control. Section 1886(d)(5)(C)(ii) of the Act requires only that "circumstances" be beyond the hospital's control. Therefore, effective with cost reporting periods beginning on or after October 1, 1987, we proposed to delete the word "extraordinary" from the regulations.

We did not receive any comments specific to our proposals for SCHs although one commenter suppported our interpretation of the law concerning the appropriateness of not paying an adjustment for volume declines in those cases in which the hospital's Medicare payment is greater than its inpatient operating costs. We note, however, that we received several comments urging changes in the criteria under which hospitals qualify as SCHs, the process for applying for SCH status, and the SCH payment methodolgy. Because these comments did not address matters presented in the proposed rule, it is not necessary that we respond to them in this final rule.

E. Referral Centers (§ 412.96)

1. Case-mix index

Section 412.96(c)(1) provides that HCFA will establish updated national and regional case-mix index values in each year's annual notice of prospective payment rates for purposes of determining referral center status. In determining the national and regional case-mix index values, we followed the same methodology we used in the November 24, 1986 final rule, as set forth in regulations at § 412.96(c)(1)(ii). Therefore, the national case-mix index value is the median case-mix index value of all urban hospitals nationwide and the regional values are the median values of urban hospitals within each census region excluding those with approved teaching programs (that is. those hospitals receiving indirect medical education payments as provided in § 412.118).

Based on bills posted to HCFA's records through February 1987 for discharges occurring during FY 1986, we proposed that to qualify for or to retain rural referral center status for cost reporting periods beginning on or after October 1, 1987, a hospital's case-mix index value for FY 1986 would have to be at least-

• 1.1594; or

· Equal to the median case-mix index value for urban hospitals (excluding hospitals with approved teaching programs as identified in § 412.118) calculated by HCFA for the census

region in which the hospital is located as indicated in the table below.

Region	Case-mix index value
1	1.1263
2	1.1136
3	1.1354
4	1.1195
5	1.0978
6	1.1492
7	1.1480
8	1.1900
9	1,1755

Based on the latest data available (discharges through June 1987), the final national case-mix index value is 1.1572 and the median case-mix index values by region are set forth in the table below.

Region	Case-mix index value
1	1.1261
2	1.1021
3	1.1355
4	1.1224
5	1.0912
6	1.1450
7	1.1442
8	1.1733
9	1.1737

We also proposed to amend § 412.96(c)(1) to state current policy that the case-mix index used to determine whether a hospital qualifies as a rural referral center is the case-mix index as calculated by HCFA from hospital billing records for Medicare discharges processed by the fiscal intermediary and submitted to HCFA's central office. This policy ensures consistency between the national and regional case-mix index standards and the case-mix index values used to determine qualification of a hospital as a rural referral center in that all case-mix index values are derived from hospitals' Medicare prospective payment bills.

For the benefit of hospitals seeking to qualify as referral centers or those wishing to know how their case-mix index value compares to the criteria, we are publishing the FY 1986 case-mix index values in Table 3c of section VI of the addendum to this rule. In keeping with our policy on discharges, these case-mix index values are computed based on all Medicare patient discharges subject to DRG-based payment. The resulting case-mix index values are based on bills received in HCFA through June 1987.

2. Discharges

Section 412.96(c)(2)(i) provides that HCFA will set forth the national and regional numbers of discharges for purposes of determining referral center status in each year's annual notice of prospective payment rates. As specified in section 1886(d)(5)(C)(i)(II) of the Act, the national standard is set at 5,000 discharges. However, we proposed to update the regional standard based on discharges for urban hospitals during the second year of the prospective payment system (that is, October 1, 1984 through September 30, 1985), which is the latest year for which we have complete discharge data available.

Therefore, we proposed that to qualify for or to retain rural referral center status for cost reporting periods beginning on or after October 1, 1987, in addition to meeting other criteria, a hospital's number of discharges for its cost reporting period that began during FY 1986 would have to be at least—

- 5,000; or
- Equal to the median number of discharges for urban hospitals in the census region in which the hospital is located as indicated in the table below.

Region	Number of dis- charges
1	6885
2	7689
3	6478
4	7848
5	6724
6	5838
7	4706
8	7157
9	4666

Based on the latest discharge data available, the final median number of discharges by census region are set forth in the table below.

Region	Number of dis- charges
1	7385
2	8192
3	6611
4	8171
5	5456
6	5879
7	4706
8	6948
9	4742

We note that there are significant differences between the proposed median regional number of discharges and the final numbers set forth above. The final discharges are based on data taken from the Health Insurance Cost Report Information System (HCRIS). The proposed median numbers were derived from data available from 2717 urban hospitals received as of April 9, 1987; the final median numbers are calculated using data from 2931 urban hospitals received as of August 20, 1987.

3. Retention Criteria

In the August 31, 1984 final rule, we stated that, once approved, a rural referral center would retain its status for three years, after which we would review the hospital's status to determine if the facility continued to meet the criteria. Basically, we stated that if the hospital met the criteria for two of the three years for which it has received the rural referral center adjustment, it would continue to qualify for payment as a rural referral center for another three-year period.

Section 9302(d)(2) of Pub. L. 99–509 extended the initial-review period for most rural referral centers by stating that, if approved on the date the law was enacted (October 21, 1986), a rural referral center would retain that status through its cost reporting period beginning before October 1, 1989. Since this would mean that the existing rural referral centers would be approved for varying lengths of time prior to their first review, we solicited comments in the proposed rule on what would be the most equitable way to evaluate these facilities for retention purposes.

4. Change in Rate Paid to Rural Referral Centers

The adjustment allowed for approved rural referral centers is that they are paid based on the urban, rather than rural, prospective payment rate as adjusted by the applicable DRG weighting factor and the rural area wage index. Section 1886(d)(5)(C)(i) of the Act provides that hospitals with approved teaching programs are not included in determining the median case mix index of urban hospitals within a census region. We define "teaching" hospitals as those hospitals receiving indirect medical education payments as provided in § 412.118.

We proposed that these same hospitals be excluded in determining the urban standardized amount paid to approved rural referral centers. We do not believe it is equitable that hospitals with approved teaching programs be excluded from the median regional case mix index calculations but be included in the calculation of the urban standardized amounts. In addition, our own analyses of Medicare cost reports from FY 1984 indicate that rural referral centers' costs, regardless of the basis upon which they qualify, are less than those of the average urban hospital when case mix, teaching status, and wage differences are taken into account. but greater than those of other rural hospitals.

We determined that deletion of the costs of urban hospitals with approved teaching programs from the calculation of the urban standardized amount would lower the amount by three percent. Therefore, instead of receiving payment based on 100 percent of the urban standardized amount, payment for approved rural referral centers would be based on 97 percent of the urban standardized amount, adjusted by the appropriate rural wage index. We proposed to amend § 412.96 (d) and (e) to implement these changes.

We received 60 letters commenting on both our proposals for rural referral centers and our current policy. One comment concerned the discharge criteria for osteopathic hospitals. Since this issue was not presented in the proposed rule, we have not responded to it in this final rule.

Comments: One commenter expressed concern regarding the process of applying the case-mix index from a past year without adjustment to determine future qualification for rural referral center status. The commenter stated that it would be more equitable to use past data to project criteria for future qualification or retention of the rural referral center adjustment.

Response: The method the commenter is suggesting is the one we used initially to establish the case-mix standards. We revised that method in our final rule published on September 3, 1986 because we believe the current method is more accurate and more equitable to hospitals.

Under section 1886(d)(5)(C)(ii) of the Act, we are required to establish the regional case-mix index standards using the median case-mix index of typical urban hospitals in each census region. We have chosen to wait until the fiscal year has ended and most claims have been processed to determine these medians. The case-mix index value of a rural hospital seeking to acquire or to

retain rural referral center status is then evaluated against these standards for the same period. This necessarily requires that the standards be determined and published for a retrospective period of time.

Basing criteria on what we believe the median case-mix standards might be could result in the disqualification of some hospitals that are similar to typical urban hospitals or result in the retention of some hospitals that have fallen below the standards. We found, in fact, that the median standards, when based on actual data, were lower than those we had published based on projections. This is demonstrated by the case-mix index standards we published in the September 3, 1986 final rule based on actual data for FY 1985 versus those in the September 3, 1985 final rule, which had been projected for the same period.

Thus, while we recognize that publishing standards based on data from prior years presents some difficulties for hospitals, we believe it is the only feasible method to ensure that rural hospitals are accurately and fairly evaluated against actual data from typical urban hospitals.

Comment: One commenter stated that while the current criteria of case-mix index, number of discharges, and medical staff composition are valid measures of a true rural referral center. considering each of these criteria individually results in some rural hospitals not qualifying for the adjustment; for example, a hospital that has a case mix index slightly below the minimum standard, but a number of discharges above the requirement together with a very high percentage of medical specialists on its staff. The commenter suggested that we create a "Referral Index" formula that would use the same criteria now in effect, but that would be formulated as follows:

Case mix index value x number of discharges + percentage of medical staff specialists [50%] = Referral Index.

Response: We believe that this is an interesting concept; however, we note that under such a concept it would be possible for a hospital to qualify as a rural referral center if it had a very low case-mix index but a high number of discharges or vice-versa.

We believe Congress intended that in order to qualify as a rural referral center, a hospital must demonstrate that both its case-mix index value and its number of discharges are comparable to a typical urban hospital in the same census region. Thus, although the suggested Referral Index might provide some relief to hospitals that narrowly miss meeting one of the criteria, but exceed the standards of one or both of

the other criteria, we believe minimum standards would still have to be imposed for each criterion.

In addition, as discussed above, current law requires that the regional criteria for both case mix index and number of discharges be based on the median of urban hospitals in the same census region. Hence, we do not have the authority to adjust the criteria with a Referral Index such as the commenter suggested.

Comment: We received identical comments from seveal hospitals suggesting that rural referral centers be paid based on the urban, rather than rural, wage index.

Response: Hospitals approved as rural referral centers have been paid based on the appropriate rural wage index since the inception of the prospective payment system. We did not propose to make any revision to this policy in our proposed rule.

We have no comments to add to our original response to this suggestion, which was presented in the January 3, 1984 final rule (49 FR 275). For the reasons cited there, we still believe the rural wage index is appropriate for rural referral centers.

Comment: One commenter suggested that the case-mix index criterion be revised to exclude teaching hospitals from the national standard as well as the regional standards. The commenter also suggested that rural referral center status be granted to a hospital if it meets the lowest standard for any census region. Finally, the commenter believes that, instead of extablishing case-mix index standards based on national and regional medians, the criterion should be based on the 25th percentile.

Response: We do not agree with any of the commenter's suggestions. We do not believe teaching hospitals should be excluded from either the national or regional case-mix index standards. (See the September 3, 1986 final rule (51 FR 31475).) However, in section 9302(d)(1) of Pub. L. 99–509, Congress required that we exclude hospitals with approved teaching programs in calculating the regional standards. Since the law did not require that teaching hospitals be excluded in calculating the national case-mix index criterion, we have not done so.

With regard to the commenters' other suggestions, section 9302(d)(1) of Pub. L. 99-509 also requires that classification of a rural hospital as a rural referral center be based (among other criteria) on whether it "has a case mix index equal to or greater than the median case

mix for hospitals * * * located in an urban area in the same region * * *."

Thus, the method for determining the regional case-mix index standards is set by law and we do not have authority to alter it.

Comment: One commenter sugested that we establish a formal appeals procedure by which a hospital seeking to acquire or to retain rural referral center status can appeal its case-mix index value.

Response: We do not believe it is necessary to establish a formal appeals process for several reasons. First, the data used to calculate each hospital's case-mix index value is taken directly from the claim forms completed by the hospital and processed by the fiscal intermediary. In section V.A. of this preamble, we have discussed the requirements and procedures to be followed if a hospital believes an incorrect DRG has been assigned to a particular claim.

Also, as discussed in our proposed notice, the same data that are used to calculate each hospital's case-mix index value are used to determine the regional and national case-mix index standards. Thus, there is consistency between and individual hospital's case-mix index value and the standard against which it

is being monitored.

Finally, since the case-mix index values published in each year's annual prospective payment update and used to determine if a hospital meets the casemix index criteria for rural referral center status are for fiscal periods that closed almost one full year prior to publication of the values, we believe the data base is virtually complete. However, should a hospital seeking rural referral status believe its published case-mix index value is inaccurate, it may request the MEDPAR data on which the case-mix index value was based. If the hospital can demonstrate that the published case-mix index value is based on an incomplete number of cases, we will recompute that hospital's case-mix index value based on Medicare bills processed by the fiscal intermediary and received in central office through the date of the hospital's application for rural referral center status.

Comment: We received numerous comments regarding our proposal to reduce the urban standardized amount used in computing payments to rural referral centers from 100 to 97 percent. All of the commenters were opposed to our proposal. Some stated that the reduction should apply only to rural referral centers that do not have approved teaching programs, that is, the commenters believe that rural referral

centers with teaching programs should continue to be paid based on 100 percent of urban standardized amount. Other commenters believe that the reduction should also be applied to all urban hospitals that do not have approved teaching programs. Still other commenters stated that the reduction should apply only to those hospitals approved under the criteria at § 412.96(c), that is, only those hospitals that qualified as rural referral centers using the case-mix index criterion.

Response: After reviewing the commenters' various objections and after further consideration, we have decided not to implement the reduction at this time. Thus, payment for all rural referral centers will continue to be based on the full 100 percent of the urban standardized amount. We will continue to study the rural referral center payment amounts and may propose adjustments in the future.

Comment: We received only two comments in response to our request for suggestions on the most equitable way to evaluate existing rural referral centers. One commenter reiterated suggestions made in the past regarding the criteria that should be used to evaluate rural referral centers and alternative methods of payment to rural referral centers. The other commenter states that our request for comments was "premature and inappropriate." Neither commenter addressed the issue of what would be an equitable time period to consider during the review process.

Response: We do not believe our solicitation of comments from interested parties on this important aspect of the rural referral center procedures was premature or inappropriate. Although all existing rural referral centers will retain their status through FY 1988, we wanted to give hospitals an opportunity to provide advice on the revised retention period criteria. We will continue to study the available options and will propose revised procedures in a future document.

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F. Payment for Services of Nonphysician Anesthetists (§ 412.113)

Section 2312 of the Deficit Reduction Act of 1984 (Pub. L. 98–369), enacted on July 18, 1984, amended sections 1886(a)(4) and 1886(d)(5) of the Act to require that we pay an additional amount to hospitals for "reasonable costs incurred" for anesthesia services furnished by certified registered nurse anesthetists (CRNAs). Section 2312(a) of Pub. L. 98–369 added section 1886(d)(5)(E) to the Act to provide for payment to hospitals on a reasonable cost basis for the costs that hospitals

incur in connection with the services of CRNAs. It further provides that this is the only payment made to the hospital for these services.

Section 1886(a)(4) of the Act, as amended by section 2312(b) of Pub. L. 98–369, exludes anesthesia services furnished by a CRNA from the definition of the term "operating costs of inpatient hospital services." Section 2312(c) of Pub. L. 98–369 specifies that these amendments are effective for hospital cost reporting periods beginning on or after October 1, 1984 and before October 1, 1987.

In implementing this provision of the law, we did not limit its application only to the services of CRNAs. The regulations at § 412.113(c) also apply the exception to the services of anesthesiology assistants. For a detailed discussion of this provision and our implementation of it, see the August 31, 1984 final rule (49 FR 34748).

Section 9320(a) of Pub. L. 99-509 amended section 2312(c) of Pub. L. 98-369 to extend the effective date of the payment on a reasonable cost basis for the services of CRNAs through cost reporting periods beginning before January 1, 1989. In the case of a cost reporting period that begins before January 1, 1989, but ends after that date, the payment made under 1886(d)(5)(E) of the Act is proportionately reduced to reflect the portion of the period occurring after January 1, 1989. Section 9320 of Pub. L. 99-509 provides that payment on a reasonable cost basis for the services of CRNAs be excluded for any part of a cost reporting period that falls after December 31, 1988. Section 9320(d) of Pub. L. 99-509 revises section 1832(a)(2)(B) of the Act to authorize direct billing for the services of CRNAs on a reasonable charge basis under Medicare Part B (Supplementary Medical Insurance) effective with services furnished on or after January 1. 1989. We proposed to revise § 412.113(c) to reflect this extension of the effective date and to make conforming changes in §§ 412.1(a), 412.2(d)(5), and 412.71(b)(8).

The Conference Committee report that accompanies Pub. L. 99–509 states that it is the intention of the conferees that the exception in § 405.553(b)(4), which permits recognition of arrangements in which physicians bill for the services of their anesthetist employees "incident to" their own services, also be extended through December 31, 1988 (H.R. 99–1012, 99th Cong., 2d Sess. 323 (1986)). (A detailed discussion of this exception is included in the September 1, 1983 interim final rule (48 FR 39794) and later revisions made to the exception are discussed in the August 31, 1984 final

rule [49 FR 34748]). We proposed, therefore, to revise § 405.553(b)(4) to reflect the extension of the exception from the usual Part B reasonable charge rules for these anesthesia services.

Comment: We received two comments on our proposal to extend the pass-through provision for nonphysician anesthetist (§ 412.113(c)) and the Part B billing exception for physicians who employ anesthetists (§ 405.553(b)(4)) until December 31, 1988. While the commenters were generally supportive, one commenter is opposed to the recognition of an arrangement that does not allow CRNAs to receive direct reimbursement independently, without the involvement of a physician.

Response: As mentioned above, the Conference Committee report that accompanied Pub. L. 99–509 clearly states Congress' intent that the Part B exception be extended for services furnished before January 1, 1989, the effective date for the direct payment of services of CRNAs on a reasonable change basis under Medicare Part B. Therefore, until that date, CRNAs cannot bill directly for their services.

VI. Other ProPAC Recommendations

As required by law, we reviewed the April 1, 1987 report submitted by ProPAC and gave its recommendations careful consideration in conjunction with the formulation of the proposals set forth in the proposed rule. We also responded to the individual recommendations in that proposed rule. The comments we received on our treatment of the ProPAC recommendations are set forth below along with our responses to those comments. However, if we received no comments from the public concerning a ProPAC recommendation or our response to that recommendation, we have not repeated the recommendation and response in the discussion below.

Recommendations 1 through 5 concerning the update factor were addressed in a separate notice that was published in the Federal Register on June 11, 1987 (52 FR 22386). Comments that were received concerning recommendations 1 through 5 are addressed in Appendix B of this document.

Recommendations 7 through 11 concerning capital were addressed in a proposed rule on that subject published in the Federal Register on May 19, 1987 (52 FR 18840). Recommendations 21, 22, and 24 through 26 concerning the DRG classification system were addressed in a proposed notice that was published in the Federal Register on May 19, 1987 (52 FR 18877).

A. Adjustments to the Payment Formula Improving the Definition of Hospital Labor Market Areas (Recommendation

Recommendation: The Secretary should adopt improved definitions of hospital labor market areas. For urban areas, the Secretary should modify the current Metropolitan Statistical Areas (MSAs) to distinguish between central and outlying areas. The central areas should be defined using urbanized areas as designated by the Census Bureau. For rural areas, the Secretary should distinguish between urbanized rural counties and other rural counties within each State. Urbanized rural counties should be defined as counties with a city or town having a population of 25,000 or greater. The implementation of improved definitions should not result in any change in aggregate hospital payments. Furthermore, these definitions should not affect the assignment of hospitals to urban or rural areas for purposes of determining standardized amounts.

Response in the Proposed Rule: For FY 1988, we do not believe that the wage index should be subdivided beyond the MSA/non-MSA distinction. Because the wage index affects every hospital's payment for every discharge, we believe additional study and analysis are necessary in order to evaluate options and determine their impact. However, as new information is developed, we will consider making improvements in labor market area definitions in future years. Our reponses regarding ProPAC's urban and rural area recommendations are as follows:

 Urban Hospitals—While subdividing urban areas into downtown "cores" and suburban "rings" could improve the explanatory power of the wage index, such subdivision would significantly increase the number of areas containing only one or two hospitals. Hospitals in these areas would enjoy a virtual pass-through of labor costs associated with Medicare hospital inpatient operating costs. Further, much of the higher wage level of core city hospitals is addressed by the teaching and disproportionate share adjustments. If we were to adopt a separate index for hospitals in urbanized areas, we would have to reconsider our policies with regard to these two adjustments.

ProPAC has recommended that urban areas be subdivided into core and ring areas on the basis of whether a hospital is located within an urbanized area. The Bureau of the Census defines an urbanized area as an area that consists of a central city or cities that, when

combined with surrounding closely settled territory ("urban fringe") having a population density of at least 1,000 persons per square mile, has a population of at least 50,000. Typically, urbanized areas cover the built-up areas at the cores of MSAs.

While we agree that the urbanized area classification may capture wage differentials, the use of urbanized areas as a basis for classifying core and ring areas may not be suitable for use in the prospective payment system. Unlike MSAs, which are county-specific, urbanized areas are defined according to actual population density and are specific to the city-block level. Also, because of the population-density basis for classifying urbanized areas, the boundaries of areas that would meet the 1,000 person per square mile criterion tend not to be stable. However, the Bureau of the Census updates urbanized areas only every 10 years. As a result, many areas that would meet the density criterion may not be classified as being in an urbanized area. Further, because urbanized areas are defined below the census-tract (and also below the MSA) level, it is not possible to determine with currently available information whether a hospital is located in an urbanized or nonurbanized area.

In summary, we do not believe that urbanized areas offer a viable system for classifying hospitals into core and ring areas because of the unstable nature of the boundaries of urbanized areas, the lag in updating urbanized areas because of the decennial census, and the inherent difficulties in determining whether a hospital is located within an urbanized area.

· Rural Hospitals-As with urban hospitals, although subdividing rural labor market areas according to urbanized and nonurbanized rural areas may increase the explanatory power of the wage index, such partition could also result in additional areas with only a few hospitals, creating for these hospitals a virtual pass through of Medicare-associated labor costs. Also, many of the high-wage rural hospitals mentioned in ProPAC's analysis are rural referral centers, which already receive the urban payment rate. In fact, analysis already indicates that large rural teaching hospitals (many of which are referral centers) are not as costly as their urban counterparts. This suggests that, even absent revisions in labor market definitions, it might be appropriate to reduce the urban rate for rural referral centers.

Further, we noted in the proposed rule that the ProPAC analysis of labor areas does not take into consideration the change in methodology required by section 9302(c) of Pub. L. 99-509, that is, computing the average standardized amounts on a discharge-weighted basis rather than on a hospital-weighted basis. We must also take into account those refinements that have already been made to the system in order to improve its equity, and how those refinements, as well as other adjustments, interact with the proposed change. For example, differential outlier offsets to the standardized rates and a reduction in the proportion of hospital costs considered to be labor-related are two changes already implemented that have increased payments to rural hospitals.

Since the factors that make up a hospital's payment are interdependent, a change in the calculation of one factor has an impact on other factors. For this reason, we believe that any analysis of redefined labor markets must be considered in the context of the payment effects to hospitals. It is not sufficient to define an improved wage index merely in terms of that index's ability to explain a greater amount of variation in hospital costs.

Further, ProPAC's recommendation does not take into account the impact of restandardization of the average costs of each hospital in the data base to reflect reconfiguration of the wage index along the lines proposed by ProPAC. In order to avoid creating overpayments and underpayments in the impact model, the same wage index, revised to reflect redefined labor market areas, must be used both in standardizing for area wage differences and in modeling payments.

In our research on the urban and rural differentials in prospective payments, we have examined the impact of alternative wage indexes and labor market areas. Overall, these alternatives produce only a marginal or modest change in prospective payments by equalizing hospital operating margins to some degree. However, it is unclear whether the redistributive effects of alternative labor market areas are appropriate. For example, an urban core-ring system would increase payments to core urban hospitals, which are generally already doing well under the prospective payment system, and decrease payments to suburban ring hospitals. Along with payment redistributions that may not be appropriate, increasing the number of labor market areas would increase the number of boundaries in the system, thereby also increasing the number of hospitals that would consider themselves unfairly disadvantaged with

respect to their location near a particular boundary.

In summary, we appreciate the work invested by ProPAC in examining labor market alternatives. However, at this point, we believe that we are still not knowledgable enough about the effects of these and other alternatives to be able to definitely recommend a particular methodology or classification system.

Comment: Several commenters expressed strong disagreement with our decision not to refine the definitions of labor market areas to encompass urban core and ring and rural urbanized and nonurbanized subdivisions. One commenter noted that, while we indicated that further study on this issue is required, the matter has been under consideration since 1985.

With regard to our contention concerning the impact of changes on individual hospitals resulting from a change in labor market areas, commenters believe that explaining variance in wage is the sole consideration in designating labor market areas. The commenters also consider invalid our argument that further subdivision of labor market areas would increase the number of hospitals enjoying a virtual pass through of wages. The commenters noted that (1) the pass-through phenomenon exists only in those subdivisions with but one hospital, (2) there is a lag between wage increases and the value of the wage index, (3) Medicare revenues represent, on average, only about 40 percent of hospital revenues, and (4) hospitals in isolated areas face market constraints on wage growth.

The commenters disagree with our contention that the instability of urbanized area boundaries is a consideration. The commenters dispute our contention that it is not possible, using currently obtained information, to determine whether hospitals are in urbanized or nonurbanized areas.

Finally, the commenters believe that rural referral center status does not address the issue that wages in urbanized rural areas are higher than wages in other rural areas.

Response: We share the commenters' concerns about the adequacy of the currend labor market area definitions. While we do not believe a revision is possible for FY 1988, we are actively reviewing refinements to labor market areas that are similar to those suggested by the commenters. Our research indicates the need to further refine urban labor market areas to reflect wage differentials across core cities, urbanized areas, and suburban areas.

Our response in the proposed rule to ProPAC's recommendation contained a lengthly explanation of our reasons for not adopting the recommendation at this time. Our concerns fall into three broad categories that we believe warrant further investigation before we propose such a revision: Impact, Methodology, and Procedure.

Impact: As noted in the proposed rule, we believe refinements to the wage index must occur in conjunction with other improvements, including refinements to the indirect medical education adjustment and disproportionate share adjustment, in order to avoid paying some hospitals or groups of hospitals too much.

Our research indicates that adoption of an urban core/ring classification based on urbanized areas would result in abrupt changes in payment, with 17 hospitals receiving less than 75 percent of their current wage index value and 226 hospitals receiving between 75 and 90 percent of their current wage index value. It should be further noted that of the 17 hospitals that would receive less than 75 percent of their current wage index value, seven had negative operating margins during the first year of the prospective payment system.

In general, adoption of an urban core/ring classification would result in many cases in which hospitals doing well would gain, and hospitals doing poorly would lose. The definitions of labor market areas are not simply based on an academic model, but rather are an integral part of a system that distributes a large amount of money nationwide. Changes to such a system ought not be taken lightly, or made simply to satisfy narrow academic criteria such as maximum explanation of variance.

In assessing the impact of adoption of a new adjustment in the prospective payment formula, one must also be cognizant of the "ripple" effect of adopting the new adjustment, that is, how other parts of the prospective payment formula are affected. If a new wage index based on redefined urban and rural areas were adopted, the labor portion of each hospital's cost per discharge in the prospective payment system data base would have to be restandardized in order to reflect the new measurement of area differences in wages. Because of this process, it is by no means the case that a hospital in an area whose wage index increases would realize increased payments under the prospective payment system. A hospital's revised Federal payment depends not only on the magnitude of any new area wage index, but also on the level of the standardized labor-

adjusted rate that is developed using that new wage index. If a hospital's wage index percentage increases less than any percentage reduction in the labor-adjusted standardized rate, then its payment will be less than under the current system. This phenomenon. which proved to be very confusing to many hospitals, was observed in the impact analysis that accompanied our report on a new wage index submitted to Congress on March 29, 1985. In view of our past experience in this matter, we believe a thorough examination of the impact of any change is necessary prior to our proposing it for adoption.

Methodology: ProPAC's recommendation is intended to better reflect hospital variation in wages. If that were the only goal, the greatest amount of wage variation could be explained by a hospital-specific wage index system. However, this is assuredly not what ProPAC is recommending, but the point at which a technical improvement ceases to be such an improvement is never specified. Certain adjustments, such as the wage index, were adopted because they explained significant amounts of variation in hospital operating costs. We find no evidence in ProPAC's report that their refined wage index provides any greater explanatory power for variation in hospital operating costs or that such improvements are sufficient to warrant the substantial administrative cost of modifying the payment system that adoption of their recommendation would entail. Moreover, as we have noted before, it is not sufficient to define an improved index only in terms of its greater explanatory power. As we stated in our first point above, the impact on payments of any revision must also be considered.

Procedure: Accepting ProPAC's recommendation would require us to adopt a whole new series of geographic definitions that have never been used in a hospital payment system. Unlike MSAs, which are familiar entities based on counties, urbanized areas are much more specific to the local level. Our data files currently do not permit us to determine a hospital's location in an urbanized area. Such location would have to be determinable with a high degree of accuracy, since hospital payment would rest on such assignment.

Comment: One commenter noted that the original reason given in the January 4, 1984 final rule (49 FR 257) for not subdividing the wage index into urban core and ring areas was that Bureau of Labor Statistics (BLS) data used in computing the wage index did not lend themselves to subdivision into small

areas. The commenter, noting that BLS data are no longer used, urged that the issue of core and ring areas be considered in the context of the overall fairness of the system.

Response: We share the commenter's concern regarding the adequacy of current labor market definitions and are studying means to further refine those definitions to more adequately reflect area wage differentials. Contrary to the commenter's assertion, there are policy considerations beyond the limitations of the BLS data. These considerations are discussed in detail in the previous response.

B. Beneficiary Concerns

1. Inpatient Hospital Cost-Sharing Requirements (Recommendation No. 18)

Recommendation: The proportion of inpatient hospital payments borne by Medicare beneficiaries should be returned to its preprospective payment system level. This proportion has inappropriately increased as a result of significant declines in length of stay experienced since the beginning of the prospective payment system. Furthermore, the structure of inpatient hospital cost-sharing requirements should be consistent with the prospective payment system incentives. In particular, current coinsurance and spell of illness requirements need to be reexamined.

Response in the Proposed Rule:
Section 9301 of Pub. L. 99–509 made a number of changes in the computation of the inpatient hospital deductible in order to make it more consistent with the current payment system. (For additional discussion of this provision, see the notice published in the Federal Register on November 20, 1986 (51 FR 42007).) In addition, the Department's recent catastrophic health proposal would further restructure the benefit package and modify beneficiary costsharing provisions.

Comment: One commenter, noting our statement that cost-sharing requirements would be restructured under the proposed catastrophic benefit legislation, urged us to reduce the amount of inpatient hospital payments borne by beneficiaries to preprospective payment levels if the legislation is not enacted.

Response: We appreciate the commenter's concern that, because of declines in the average length of stay, the inpatient deductible and coinsurance requirement have risen to a higher percent of the average cost of a hospital stay. However, as noted above, we believe that this problem has been resolved by the changes made by

section 9301 of Pub. L. 99–509 and by the fact that recent evidence indicates that the average length of stay is no longer decreasing. Nevertheless, we are reviewing the Medicare cost-sharing requirements, and if the catastrophic health insurance legislation is not enacted, we will explore ways to mitigate the amount of payment made by beneficiaries, consistently with budgetary concerns.

2. Evaluating the Results of PRO Quality of Care Review (Recommendation No. 19)

Recommendation: The Secretary should promptly initiate a comprehensive evaluation of PRO quality of care review activities and findings. The evaluation should assess the impact on quality of care of preadmission, admission, transfer, and readmission review activities. The PRO findings concerning quality of the services furnished during an admission and the health outcome of the episode of care should also be evaluated. ProPAC is aware that the Super-PRO is auditing and validating PRO review activities. However, ProPAC does not believe that this effort can substitute for a comprehensive evaluation of the extent to which PROs are identifying, assessing, and correcting problems related to quality of care.

Response in the Proposed Rule: We have an extensive and comprehensive system in place to evaluate the credibility of PRO review decisions. including those related to quality of care. ProPAC does not consider the "Super-PRO" evaluation of PRO medical determinations to be sufficient to monitor PRO findings. We agree that the "Super-PRO" alone is not sufficient. However, if the "Super-PRO" results are viewed in the context of other evaluation activities, we believe that we are adequately assessing PRO performance in the area of quality of care review. We believe ProPAC's recommendation would result in a duplicative evaluation effort.

Comment: One commenter was concerned that use of the "Super-PRO" to audit and validate PRO review activities was not a substitute for a comprehensive review and evaluation of the PROs to determine the extent to which PROs are identifying, assessing, and correcting problems related to quality of care.

Response: We agree with the commenter's concern that the impact of PRO review on the patterns of quality of care should be the focus of a substantial evaluation and we have begun such a process. Section 9353(c)(3) of Pub. L. 99—

509 requires that we identify methods that are available to PROs to identify cases that are more likely than others to be associated with substandard quality of care. These methods could include statistical profiling (now required of all PROs) of practice patterns of physicians and providers and of DRG assignments, and subsequent identification of areas on which to focus review. We have provided funds to several PROs to develop a clinical data base and, eventually, a means of screening out cases (or patterns of cases) that are not likely to be quality of care problems. The data produced by these two activities can then be used to compare patterns of care both before and after PRO intervention.

In addition, the 1986–1988 PRO Scope of Work requires that PROs use definitive quality of care methodologies, including generic quality screens. Our data analysis at the end of the current contract period should enable us to evaluate the impact of those screens in terms of confirmation of quality of care problems, to determine whether patterns exist, and to establish whether trends are present.

PRO performance will be comprehensively evaluated by us and the results will be used in making decisions or contract renewals. We are developing a procedure to release these data to the public.

C. DRG Classification and Weighting Factors

Additional Payment for Magnetic Resonance Imaging (MRI) Scans (Recommendation No. 27)

Recommendation: For a three-year period, Medicare should pay hospitals an additional amount to reflect operating costs for each covered magnetic resonance imaging (MRI) scan performed on an inpatient Medicare beneficiary in a prospective payment hospital. The add-on payment should be calculated by the Secretary each year to reflect both changes in the average cost of an efficiently produced scan and the degree to which MRI substitutes for other hospital procedures.

Response in the Proposed Rule: We recognize ProPAC's concern that the current payment methodology may act as a disincentive to the widespread use of MRI technology. However, we regard this concern as anticipatory, since there is no evidence that hospitals furnishing MRI are losing money under the prospective payment system. On the contrary, the hospitals most likely to be furnishing MRI services are urban teaching hospitals; that is, the institutions that have been faring the

best under the prospective payment system. We have always held that one of the basic tenets of a system built on averages is that payments would not cover costs in all cases and that excess payments on some cases would offset losses in other cases.

We are concerned that there will be numerous technological advances in the future that would be similar to MRI; that is, several DRGs would be affected by the changes. If we begin to unbundle the prospective payment rate to provide add-on payments in that manner, the basic concept of prospective payments on a discharge basis would be undermined.

We are, however, giving the issue further study. Unique ICD-9-CM codes for MRI services were approved effective October 1, 1986. From these data, we will be able to evaluate the issue more thoroughly in the upcoming months. If we find that the current prospective payment methodology adversely affects the quality of care, we will consider alternative payment options, including add-ons.

Comment: Several commenters supported the concept of an add-on payment for MRI cases. The commenters noted that this type of payment is necessary to encourage diffusion of this new technology and to ensure that MRI scans are made available to those beneficiaries who need them.

Response: As we stated in the proposed rule, although we are not providing an add-on payment for MRI, we have approved unique ICD-9-CM codes for MRI procedures in order to evaluate the adequacy of payment. We believe that add-on payments for new technology, including MRI, should be made in only the most compelling cases. Approval of an add-on for one technology would set a precedent that would encourage subsequent petitions for additional payments for other technologies. Further, as we discussed in the proposed rule, add-ons for MRI would undermine one of the basic tenets of the prospective payment system; that is, that the payment level made for each case is based on an average. Finally, it is not the function of the prospective payment system either to encourage or to discourage the diffusion of new technologies.

VII. Other Required Information

A. Effective Dates

The effective date of this final rule (including the addendum and appendixes) is October 1, 1987. The following change is effective beginning with cost reporting periods beginning on or after October 1, 1987: Section

412.92(e)(2)(ii)—Special treatment: Sole community hospitals.

B. Paperwork Reduction Act

This final rule does not impose information collection requirements. Consequently, it need not be reviewed by the Executive Office of Management and Budget under the authority of the Paperwork Reduction Act of 1980 (44 U.S.C. 3501–3511).

List of Subjects

42 CFR Part 405

Administrative practice and procedure, Health facilities, Health professions, Kidney diseases, Laboratories, Medicare, Nursing homes, Reporting and recordkeeping requirements, Rural areas, X-rays.

42 CFR Part 412

Health facilities, Medicare.

42 CFR Part 413

Administrative practice and procedure, Health facilities, Health professions, Kidney diseases, Laboratories, Medicare, Nursing homes, Reporting and recordkeeping requirements, Rural areas, X-rays.

42 CFR Part 466

Competitive medical plans (CMPs), Grant programs—health, Health care, Health facilities, Health maintenance organizations (HMOs), Health professions, Peer Review Organizations.

42 CFR Chapter IV is amended as follows:

Chapter IV—Health Care Financing Administration

Department of Health and Human Services Subchapter B-Medicare Programs

I. Part 405, Subpart E is amended as follows:

PART 405—FEDERAL HEALTH INSURANCE FOR THE AGED AND DISABLED

Subpart E—Criteria for Determination of Reasonable Charges; Reimbursement for Services of Hospital Interns, Residents, and Supervising Physicians

A. The authority citation for Subpart E continues to read as follows:

Authority: Secs. 1102, 1814(b), 1832, 1833(a), 1842(b) and (h), 1861(b) and (v), 1862(a)(14), 1866(a), 1871, 1881, 1886, 1887, and 1889 of the Social Security Act as amended (42 U.S.C. 1302, 1395f(b), 1395k, 1395l(a), 1395u(b) and (h), 1395x(b) and (v), 1395y(a)(14), 1395cc(a), 1395hh, 1395rr, 1395ww, 1395xx, and 1395zz).

§ 405.553 [Amended]

B. In § 405.553, in paragraph (b)(4), the phrase "a cost reporting period beginning on or after October 1, 1984 and before October 1, 1987." is revised to read "cost reporting periods beginning on or after October 1, 1984 through any part of a cost reporting period occurring before January 1, 1989."
II. Part 412 is amended as follows:

PART 412—PROSPECTIVE PAYMENT SYSTEM FOR INPATIENT HOSPITAL SERVICES

A. The authority citation for Part 412 continues to read as follows:

Authority: Secs. 1102, 1122, 1871, and 1886 of the Social Security Act, as amended (42 U.S.C. 1302, 1320a-1, 1395hh, and 1395ww)

B. Subpart A is amended as follows:

Subpart A-General Provisions

§ 412.1 [Amended]

1. a. In § 412.1(a), in the third sentence, the phrase "and before October 1, 1987," is revised to read "through any part of a cost reporting period occurring before January 1, 1989."

b. In § 412.1(b), a new sentence is added at the end of the paragraph to read "Subpart K describes how the prospective payment system is implemented for hospitals located in Puerto Rico."

§ 412.2 [Amended]

2. In § 412.2(d)(5), the phrase "and before October 1, 1987," is revised to read "through any part of a cost reporting period occurring before January 1, 1989,"

C. In Subpart B, § 412.23(f) is revised to read as follows:

Subpart B-Hospital Services Subject to and Excluded from the Propective **Payment System**

§ 412.23 Excluded hospitals: Classifications.

(f) Hospitals outside the 50 States, the District of Columbia, or Puerto Rico. A hospital is excluded from the prospective payment system if it is not located in one of the fifty States, the District of Columbia, or Puerto Rico. * * * *

D. Subpart D is amended as follows:

Subpart D-Basic Methodology for **Determining Federal Prospective Payment Rates**

1. In § 412.60, paragraph (d) is redesignated as paragraph (e), a new paragraph (d) is added, and newly

redesignated paragraph (e) is revised to read as follows:

§ 412.60 DRG classification and weighting factors.

(d) Review of DRG assignment. (1) A hospital has 60 days after the date of the notice of the initial assignment of a discharge to a DRG to request a review of that assignment. The hospital may submit additional information as a part of its request.

(2) The intermediary reviews the hospital's request and any additional information and decides whether a change in the DRG assignment is appropriate. If the intermediary decides that a higher-weighted DRG should be assigned, it must request the appropriate PRO to review the case to verify the change in DRG assignment as specified in § 466.70(e)(2) of this chapter.

(3) Following the 60-day period described in paragraph (d)(1) of this section, the hospital may not submit additional information with respect to the DRG assignment or otherwise revise

(e) Revision of DRG classification and weighting factors. Beginning with discharges in fiscal year 1988, HCFA adjusts the classifications and weighting factors established under paragraphs (a) and (b) of this section at least annually to reflect changes in treatment patterns, technology, and other factors that may change the relative use of hospital resources.

2. In § 412.63, text is added to paragraph (f) to read as follows:

§ 412.63 Federal rates for fiscal years after Federal fiscal year 1984.

(f) Applicable percentage change for fiscal year 1988. The applicable percentage change for fiscal year 1988 is the percentage increase in the market basket index (as described in § 413.40(c)(3)(ii) minus 2.0 percentage points.

E. Subpart E is amended as follows:

Subpart E-Determination of **Transition Period Payment Rates**

§ 412.71 [Amended]

1. In § 412.71(b)(8), the phrase "October 1, 1984, and before October 1, 1987," is revised to read "on or after October 1, 1984 through any part of a cost reporting period occurring before January 1, 1989,".

2. In § 412.73, text is added to a new paragraph (c)(5) and reserved paragraph (c)(6) is removed to read as follows:

§ 412.73 Determination of the hospitalspecific rate.

(c) Updating base-year costs. *

(5) For Federal fiscal year 1988 and following. For purposes of determining the prospective payment rates for sole community hospitals under § 412.92(d). the base-year cost per discharge continues to be updated each Federal fiscal year as follows:

(i) For Federal fiscal year 1988, the update factor is the percentage increase in the market basket index (as described in § 413.40(c)(3)(ii)) minus 2.0 percentage

points.

(ii) For Federal fiscal years 1989 and following, the update factor is determined using the methodology set forth in § 412.63(g)(1) through (g)(3). * * *

F. Subpart G is amended as follows:

Subpart G-Special Treatment of **Certain Facilities**

1. In § 412.92, the introductory text of paragraph (e)(2) is republished and paragraph (e)(2)(ii), the introductory language of paragraph (e)(3), and paragraph (e)(3)(i) are revised to read as follows:

§ 412.92 Special treatment: Sole community hospitals.

(e) Additional payments to sole community hospitals experiencing a significant volume decrease during the transition period. *

(2) To qualify for a payment adjustment on the basis of a decrease in discharges, a sole community hospital

(ii) Show that the decrease is due to circumstances beyond the hospital's

(3) HCFA determines a lump sum adjustment amount not to exceed the difference between the hospital's Medicare inpatient operating costs and the hospital's total DRG revenue based on DRG-adjusted prospective payment rates (including outlier payments determined under Subpart F of this part and additional payments made for hospitals that serve a disproportionate share of low-income patients as determined under § 412.106 and for indirect medical education costs as determined under § 412.118). In determining the adjustment amount, HCFA considers-

(i) The individual hospital's needs and circumstances, including the reasonable cost of maintaining necessary core staff

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and services in view of minimum staffing requirements imposed by State agencies:

2. In § 412.96(c)(1), the introductory language is revised to read as follows:

§ 412.96 Special treatment: Referral centers.

(c) Alternative criteria.* * *

(1) Case-mix index. HCFA sets forth national and regional case-mix index values in each year's annual notice of prospective payment rates published under § 412.8(b). The methodology HCFA uses to calculate these criteria is described in paragraph (g) of this section. The case-mix index value to be used for an individual hospital in the determination of whether it meets the case-mix index criteria is that calculated by HCFA from the hospital's own billing records for medicare discharges as processed by the fiscal intermediary and submitted to HCFA. The hospital's casemix index for discharges (not including discharges from distinct part units excluded from the prospective payment system under Subpart B of this part) during the Federal fiscal year that ended one year prior to the beginning of the cost reporting period for which the hospital is seeking referral center status must be at least equal to-

G. In Subpart H, § 412.113 is amended as follows:

Subpart H—Payments to Hospitals under the Prospective Payment System

§ 412.113 [Amended]

In § 412.113(c), the phrase "and before October 1, 1987," is revised to read "through any part of the cost reporting period occurring before January 1, 1989,".

H. A new Subpart K is added to read as follows:

Subpart K—Prospective Payment System for Hospitals Located in Puerto Rico

Sec

412.200 General provisions.

412.204 Payments to hospitals located in Puerto Rico.

412.208 Puerto Rico rates for Federal fiscal year 1988.

412.210 Puerto Rico rates for fiscal years after Federal fiscal year 1988.

412.212 National rate.

412.220 Special treatment of certain hospitals located in Puerto Rico.

Subpart K—Prospective Payment System for Hospitals Located in Puerto Rico

§ 412.200 General provisions.

Beginning with discharges occurring on or after October 1, 1987, hospitals located in Puerto Rico are subject to the rules governing the prospective payment system. Except as provided in this subpart, the provisions of Subparts A. B. C, F, G, and H of this part apply to hospitals located in Puerto Rico, Except for § 412.60, which deals with DRG classification and weighting factors, the provisions of Subpart D and E, which describe the methodology used to determine prospective payment rates for hospitals, do not apply to hospitals located in Puerto Rico. Instead, the methodology for determining porspective payment rates for these hospitals is set forth in §§ 412.204 through 412.212.

§ 412.204 Payments to hospitals located in Puerto Rico.

Payments to hospitals located in Puerto Rico that are paid under the prospective payment system are equal to the sum of —

(a) 75 percent of the Puerto Rico prospective payment rate, as determined under § 412.208 or § 412.210; and

(b) 25 percent of a national prospective payment rate, as determined under § 412.212.

§ 412.208 Puerto Rico rates for Federal fiscal year 1988.

(a) General rule. HCFA determines the Puerto Rico adjusted DRG prospective payment rate for each inpatient hospital discharge occurring in Federal fiscal year 1988 for a prospective payment hospital. These rates are determined as described in paragraphs (b) through (i) of this section.

(b) Determining target amounts. For each hospital subject to the prospective payment system. HCFA determines the Medicare target amount, as described in § 413.40(c) of this chapter, for the hospital's cost reporting period beginning in fiscal year 1967. Revisions in the target amounts made subsequent to establishment of the standardized amounts under paragraph (d) of this section do not affect the standardized amounts.

(c) Updating the target amounts for fiscal year 1988. HCFA updates each target amount determined under paragraph (b) of this section for fiscal year 1988 by prorating the applicable percentage increase (as defined in § 412.63(f) of this chapter) for fiscal year 1988 to the midpoint of fiscal year 1988 (April 1, 1988).

(d) Standardizing amounts. HCFA standardizes the amount updated under paragraph (c) of this section for each hospital by—

(1) Adjusting for variations in case mix among hospitals;

(2) Excluding an estimate of indirect medical education costs;

(3) Adjusting for area variations in hospital wage levels; and

(4) Excluding an estimate of the payments for hospitals that serve a disproportionate share of low-income patients.

(e) Computing urban and rural averages. HCFA computes separate discharge-weighted averages of the standardized amounts determined under paragraph (d) of this section for urban and rural hospitals in Puerto Rico.

(f) Geographic classifications. (1) For purposes of paragraph (e) of this section, the following definitions apply:

(i) The term "urban area" means a Metropolitan Statistical Area (MAS), as defined by the Executive Office of Management and Budget.

(ii) The term "rural area" means any area outside an urban area.

(2) A hospital classified as rural is deemed to be urban and receives the urban Puerto Rico payment amount if the county in which it is located meets the following criteria:

(i) At least 95 percent of the perimeter of the rural county is contiguous with urban counties.

(ii) The county was reclassified from an urban area to a rural area after April 20, 1983, as described in § 412.62(f)(1)(iv).

(iii) At least 15 percent of employed workers in the county commute to the central county of one of the adjacent MSAs.

(g) Reducing for value of outlier payments. HCFA reduces each of the average standardized amounts determined under paragraphs (c) through (e) of this section by a proportion equal to the proportion (estimated by HCFA) of the total amount of payments based on DRG prospective payment rates that are additional payments to hospitals located in Puerto Rico for outlier cases under Subpart F of this part.

(h) Computing Puerto Rico rates for urban and rural hospitals. For each discharge classified within a DRG, HCFA establishes a Puerto Rico prospective payment rate, as follows:

(1) For hospitals located in an urban area, the rate equals the product of—

(i) The average standardized amount (computed under paragraphs (c) through (g) of this section) for hospitals located in an urban area; and (ii) The weighting factor determined under § 412.60(b) for that DRG.

(2) For hospitals located in a rural area, the rate equals the product of—

(i) The average standardized amount (computed under paragraphs (c) through (g) of this section) for hospitals located in an urban area; and

(ii) The weighting factor determined under § 412.60(b) for that DRG.

(i) Adjusting for different area wage levels. HCFA adjusts the proportion (as estimated by HCFA from time to time) of Puerto Rico rates computed under paragraph (h) of this section that are attributable to wages and labor-related costs, for area differences in hospital wage levels, by a factor (established by HCFA) reflecting the relative hospital wage level in the geographic area of the hospital compared to the national average hospital wage level.

§ 412.210 Puerto Rico rates for fiscal years after Federal fiscal year 1988.

(a) General rule. (1) HCFA determines the Puerto Rico adjusted prospective payment rate for each inpatient hospital discharge occurring in a Federal fiscal year after fiscal year 1988 that involves inpatient hospital services of a hospital in Puerto Rico subject to the prospective payment system for which payment may be made under Medicare Part A.

(2) The rate is determined for hospitals located in urban or rural areas within Puerto Rico, as described in paragraphs (b) through (e) of this

section.

(b) Geographic classifications. For purposes of this section, the definitions set forth in § 412.208(f) apply.

(c) Updating previous standardized amounts. HCFA computes separate average standardized amounts for hospitals in urban areas and rural areas within Puerto Rico equal to the respective average standardized amount computed for fiscal year 1988 under § 412.208(e)—

(1) Increased by the applicable percentage change determined under

§ 412.63(g); and

(2) Reduced by a proportion equal to the proportion (estimated by HCFA) of the total amount of prospective payments that are additional payment amounts to hospitals located in Puerto Rico attributable to outlier cases under Subpart F of this part.

(d) Computing Puerto Rico rates for

(d) Computing Puerto Rico rates for urban and rural hospitals. For each discharge classified within a DRG, HCFA establishes for the fiscal year a Puerto Rico prospective payment rate as

follows:

(1) For hospitals located in an urban area in Puerto Rico, the rate equals the product of—

(i) The average standardize amount (computed under paragraph (c) of this section) for the fiscal year for hospitals located in an urban area; and

(ii) The weighting factor determined under § 412.60(b) for that DRG.

(2) For hospitals located in a rural area in Puerto Rico, the rate equals the product of—

(i) The average standardized amount (computed under paragraph (c) of this section) for the fiscal year for hospitals located in a rural area; and

(ii) The weighting factor (determined under § 412.60(b)) for that DRG.

(e) Adjusting for different area wage levels. HCFA adjusts the proportion (as estimated by HCFA from time to time) of Puerto Rico rates computed under paragraph (d) of this section that is attributable to wages and labor-related costs for area differences in hospital wage levels by a factor (established by HCFA) reflecting the relative hospital wage level in the geographic area of the hospital compared to the national average hospital wage level.

§ 412.212 National rate.

(a) General rule. For purposes of payment to hospitals located in Puerto Rico, the national prospective payment rate is determined as described in paragraphs (b) through (d) of this section.

(b) Computing a national average standardized amount. HCFA computes a discharge-weighted average of the—

 National urban adjusted standardized amount determined under § 412.63(i)(1)(i); and

(2) National rural adjusted average standardized amount determined under § 412.63(i)(2)(i).

(c) Computing a national rate. For each discharge classified within a DRG, the national rate equals the product of—

 The national average standardized amount computed under paragraph (b) of this section; and

(2) The weighting factor (determined under § 412.60(b)) for that DRG.

(d) Adjusting for different area wage levels. HCFA adjusts the proportion (as estimated by HCFA from time to time) of the national rate computed under paragraph (c) of this section that is attributable to wages and labor-related costs for area differences in hospital wage levels by a factor (established by HCFA) reflecting the relative hospital wage level in the geographic area of the hospital compared to the national average hospital wage level.

§ 412.220 Special treatment of certain hospitals located in Puerto Rico.

Subpart G of this part sets forth rules for special treament of certain facilities

under the prospective payment system.
The following sections in Subpart G of
this part do not apply to hospitals
located in Puerto Rico:

(a) Section 412.92, sole community hospitals.

(b) Section 412.96, referral centers.

III. Part 413 is amended as follows:

PART 413—PRINCIPLES OF REASONABLE COST REIMBURSEMENT; PAYMENT FOR END-STAGE RENAL DISEASE SERVICES

A. The authority citation for Part 413 continues to read as follows:

Authority: Sections 1102, 1122, 1814(b), 1815, 1833(a), 1861(v), 1871, 1881, and 1886 of the Social Security Act as amended (42 U.S.C. 1302, 1320a-1, 1395f(b), 1395g, 1395f(a), 1395x(v), 1395hh, 1395rr, and 1395ww).

B. In Subpart C, § 413.40, the introductory text in paragraph (c)(3)(i) is republished and text is added to paragraph (c)(3)(i)(C) to read as follows:

§ 413.40 Ceiling on rate of hospital costs increases.

(c) Procedure for establishing the ceiling (target amount).

(3) Target rate percentage.

(i) The applicable target rate percentage is determined as follows:

(C) Federal fiscal year 1988. The applicable target rate percentage for cost reporting periods beginning on or after October 1, 1987 and before October 1, 1988 is the percentage increase in the market basket index minus 2.0 percentage points.

IV. Part 466, Subpart C is amended as follows:

PART 466—UTILIZATION AND QUALITY CONTROL REVIEW

Subpart C—Review Responsibilities of Utilization and Quality Control Peer Review Organizations (PROs)

A. The authority citation for Part 466 continues to read as follows:

Authority: Secs. 1102, 1154, and 1871 of the Social Security Act (42 U.S.C. 1302, 1320c-3, and 1395hh).

B. In § 466.70, paragraph (e) is amended by redesignating paragraph (e)(2) as (e)(3) and adding a new paragraph (e)(2) to read as follows:

§ 466.70 Statutory bases, applicability and provisions.

(e) Other duties and functions.

(2) The PRO must review every change in a DRG assignment that is a result of a review made under the provisions of § 412.60(d) if the change results in the assignment of a higher-weighted DRG and the PRO has not previously reviewed the case. The PRO must verify that the diagnostic and procedural information supplied by the hospital is substantiated by the information in the medical record.

(Catalog of Federal Domestic Assistance Program No. 13.773, Medicare—Hospital Insurance Program)

Dated: August 24, 1987.

William L. Roper,

Administrator, Health Care Financing Administration.

Approved: August 26, 1987.

Otis R. Bowen,

Secretary.

[Editorial Note.—The following addendum and appendixes will not appear in the Code of Federal Regulations.]

Addendum—Schedule of Standardized Amounts Effective With Discharges On or After October 1, 1987, and Update Factors and Target Rate Percentages Effective With Cost Reporting Periods Beginning On or After October 1, 1987

1. Summary and Background

In this addendum to the final rule, we are making changes in the methods, amounts, and factors for determining prospective payment rates for Medicare inpatient hospital services. We are also setting forth the methods, amounts, and factors for determining prospective payment rates for Medicare inpatient hospital services furnished by hospitals in Puerto Rico. Finally, we are setting new target rate percentages for determining the rate-of-increase limits (target amounts) for hospitals and hospital units excluded from the prospective payment system.

For hospital cost reporting periods beginning on or after October 1, 1987, except for sole community hospitals and hospitals located in Puerto Rico, each hospital's payment per discharge under the prospective payment system will, for the first time, be comprised of 100 percent of the Federal rate; that is, hospitals will no longer receive any part of their payment based on a hospitalspecific rate (section 1886(d)(1)(A) of the Act). That section of the Act also requires that for discharges occurring on or after October 1, 1987, the Federal portion of a hospital's prospective payment rate is based on 100 percent of the national rate, instead of a blend of regional and national rates.

Sole community hospitals will continue to be paid on the basis of a rate per discharge composed of 75 percent of the hospital-specific rate and 25 percent of the applicable Federal regional rate (section 1886(d)(5)(C)(ii) of the Act).

Effective with discharges occurring on or after October 1, 1987, hospitals in Puerto Rico will be subject to the prospective payment system (section 1886(d)(9) of the Act as added by section 9304(a) of Pub. L. 99–509). However, these hospitals' payment per discharge will be the sum of 75 percent of a Puerto Rico rate and 25 percent of a national rate.

As discussed below in section II, we are making changes in the determination of the prospective payment rates. The changes, to be applied prospectively, will affect the calculation of the Federal rates. Section III sets forth our determination of payment rates for hospitals in Puerto Rico. In section IV. we discuss the various adjustments made to the average standardized amounts in order to achieve budget neutrality in those areas in which it is required. Section V sets forth our changes for determining the rate-ofincrease limits for hospitals excluded from the prospective payment system. The tables to which we refer in the preamble to the final rule are presented at the end of this addendum.

II. Changes to Prospective Payment Rates and DRG Weighting Factors for FY 1988

The basic methodology for determining Federal national prospective payment rates is set forth at § 412.63. Below, we discuss the manner in which we are changing some of the factors or methodologies used for determining the prospective payment rates. The Federal rate changes will be effective with discharges occurring on or after October 1, 1987.

In summary, we are establishing the FY 1988 national and regional rates (that is, the standardized amounts set forth in Table 1a and 1b of the addendum) by—

Restandardizing, with the 1982
 HCFA wage index, the hospital costs used to establish the rates to reflect the revisions we are making in the methodology for calculating the national average hourly wage;

 Computing average costs per case per hospital and adjusting costs per case to exclude the effects of case mix, indirect medical education costs, payment adjustments to disproportionate share hospitals, and cost-of-living differences for Alaska and

 Grouping the adjusted operating costs per case (labor-related and nonlabor-related) to compute urban and rural, national and regional average standardized amounts using averages weighted by total discharges rather than by number of hospitals;

 Updating the standardized amounts by 2.7 percent (that is, the increase in the market basket percentage minus 2.0

percentage points).

A. Calculation of Adjusted Standardized Amounts

1. Standardization and Restandardization of Base-Year Costs. Section 1886(d)(2)(A) of the Act required the establishment of base-year cost data containing allowable operating costs per discharge of inpatient hospital services for each hospital. The preamble to the interim final rule, published September 1, 1983 (48 FR 39763), contains a detailed explanation of how base-year cost data were established in the initial development of standard amounts for the prospective payment system and how they are used in computing the Federal rates.

Section 1886(d)(2)(C) of the Act required that the updated base-year per discharge costs be standardized in order to remove from the cost data the effects of certain sources of variation in cost among hospitals. These include case mix, differences in area wage levels, cost of living adjustments, and indirect medical education costs. We proposed to restandardize the base-year costs using the 1982 HCFA wage index to reflect the change in the methodology for computing the national average hourly wage.

We did not propose to restandardize the base-year costs for the following:

· Case mix.

Indirect medical education costs.

• Cost of living for Alaska and Hawaii.

 Payments to hospitals that serve a disproportionate share of low-income patients.

a. Adjustments for Variation in Hospital Wage Levels. Section 1886(d)(2)(C)(ii) of the Act requires that we standardize the average cost per case of each hospital used to develop the separate urban and rural standardized amounts for differences in area wage levels. Therefore, we divided each standardized amount into labor and nonlabor portions, based on the labor and nonlabor components of the hospital market basket, and standardized the labor portion of the FY 1984 standardized amounts using the Bureau of Labor Statistics (BLS's) area wage index. For FY 1986, we adopted a new wage index based on HCFA survey data, and we restandardized the base

year costs used to calculate the FY 1986 standardized amounts to account for the new wage index. We removed the effect of the previous standardization for each hospital's BLS wage index by multiplying each hospital's average cost per discharge value by the old index and restandardized the amounts by dividing that result by the new HCFA wage index (see 50 FR 35692).

As discussed in section III of the preamble, we proposed to use a blended HCFA wage index composed of two separate wage indexes based on 1982 and 1984 data, respectively, and to make a change in the methodology for computing the national average hourly wage, which serves as the basis for indexing the area wage levels. However, the latter change would result in lower index values for all areas relative to the national average hourly wage, since the national average hourly wage based on the 1982 data is higher using the proposed methodology. In order for our porposed change in methodology to have no adverse impact on level of payments to hospitals, the base year costs used to calculate the standardized amounts must be restandardized to take into account the effect on each area's wage index value of the revised methodology for calculating the national average hourly wage.

Therefore, we proposed to restandardize the base year costs that were used to calculate the standardized amounts using the 1982 HCFA wage index. We removed the effect of the previous standardization (1982 HCFA wage index based on an area-weighted national average hourly wage) by multiplying each hospital's average cost per discharge value by the current 1982 wage index and restandardizing the amount by dividing that result by the 1982 HCFA wage index recalculated using the proposed methodology for computing the national average hourly

b. Variations in Case Mix Among Hospitals. Section 1886(d)(2)(C)(iii) of the Act requires that the updated FY 1984 amounts be standardized to adjust for variations in case mix among hospitals. The methodology used for determining the appropriate adjustment factor (that is, the case-mix index) is explained in the September 1, 1983 interim final rule (48 FR 39768-39771). A case-mix index has been calculated for

each hospital based on 1981 cost and billing data.

Standardization, necessary to neutralize inpatient operating costs for the effects of variations in case mix, is accomplished by dividing the hospital's average cost per Medicare discharge by that hospital's case-mix index. Table 3a in the addendum to the September 1. 1983 interim final rule (48 FR 39847-39870) contains the case-mix index values used for this purpose. We did not propose to make any changes to the case-mix index for inpatient operating costs and, therefore, did not restandardize the updated amounts for variations in case mix.

c. Indirect Medical Education Costs. Section 1886[d][2][C][i] of the Act requires that the updated FY 1984 amounts be standardized for indirect medical education costs. Section 1886(d)(5)(B) of the Act provides that prospective payment hospitals receive an additional payment for the indirect costs of medical education. Section 9104(a) of Pub. L. 99-272 revised section

1886(d)(5)(B) of the Act to change the education adjustment factor used to determine the indirect medical education payment. Section 1886(d)(5)(B) of the Act currently specifies that the education adjustment factor is approximately 8.1 percent for discharges occurring on or after May 1, 1986 and before October 1, 1989. For discharges occurring on or after October 1, 1989, the adjustment factor is equal to approximately 8.7 percent. These factors are approximations because they are applied on a curvilinear or variable basis, rather than on a linear basis. An adjustment made on a curvilinear basis reflects a nonlinear cost relationship. that is, each absolute increment in a hospital's ratio of interns and residents to beds does not result in an equal proportional increase in costs. Therefore, the adjustment factors are only approximately 8.1 percent and 8.7 percent.

For discharges occurring on or after May 1, 1986 and before October 1, 1989, the indirect medical education factor

equals the following:

For discharges occurring on or after October 1, 1989, the indirect medical education factor equals the following:

Section 9104(b) of Pub. L. 99-272 amended section 1886(d)(2)(C)(i) of the Act to provide that the standardized amounts be restandardized to reflect the changes made to the payment adjustment for indirect medical education costs under section 9104(a) of Pub. L. 99-272. Therefore, in establishing the standardized amounts used to determine the FY 1987 prospective payment rates, after adjusting each hospital's inpatient operating cost per discharge for inflation, differences in area wage levels, and case mix, we divided each teaching hospital's cost per discharge by 1.0 plus the individual hospital's indirect medical education adjustment factor as computed using the formula described above, which section 1886(d)(5)(B)(ii)(I) of the Act requires be used for discharges on or after May 1,

1986 and before October 1, 1989. We did not propose to restandardize the baseyear costs for FY 1988 for indirect medical education costs.

d. Cost-of-Living Factor for Alaska and Hawaii. Section 1886(d)(5)(C)(iv) of the Act authorizes the Secretary to provide for such adjustments to the payment amounts as the Secretary deems appropriate to take into account the unique circumstances of hospitals located in Alaska and Hawaii.

Generally, these two States have higher levels of cost in comparison to other States in the nation. The high cost of labor is accounted for in the wage index adjustments discussed above. However, the high cost of living in these States also affects the cost of nonlabor items (for example, supplies and

equipment). Therefore, in order to remove the effects of the higher nonlabor costs from the overall cost data (that is, for standardization purposes), the nonlabor portion of the average cost per Medicare discharge in hospitals located in Alaska and Hawaii is divided by an appropriate cost-of-living adjustment factor.

e. Costs for Hospitals that Serve a Disproportionate Share of Low-Income Patients. Section 1886(d)(2)(C)(iv) of the Act provides that, effective with discharges occurring on or after October 1, 1986 and before October 1, 1989, the updated amounts be standardized for the estimated additional payments made to hospitals that serve disproportionate shares of low-income patients. That is, the law requires us to remove the effects of the payments made to disproportionate share hospitals from the costs used to establish the standardized amounts. For discharges occurring on or after October 1, 1989, we will no longer make such an adjustment to take into account the estimated payments made to disproportionate share hospitals, since section 1886(d)(5)(F) of the Act does not authorize such payments for discharges after September 30, 1989.

Therefore, in establishing the standardized amounts for FY 1988, we proposed to adjust each disproportionate share hospital's inpatient operating cost per discharge by adding 1.0 to the applicable disproportionate share payment factor, and dividing the hospital's cost per discharge by that number. In this way, we removed the effect of payment adjustments for disproportionate share hospitals from the standardized amounts as required under section 1886(d)(2)(C)(iv) of the Act.

Under section 1886(d)(5)(F)(iv) calculation of the disproportionate share adjustment factor requires us to calculate the number of a hospital's patient days attributable to Medicare beneficiaries entitled to Supplemental Security Income (SSI), and to non-Medicare beneficiaries eligible for Medicaid. In determining the disproportionate share adjustment factors for purposes of standardizing the standardized amounts, we proposed to use available data on the percentage of Medicaid days from Medicare cost reports with cost reporting periods beginning in Federal FY 1984, and we proposed to use the percentage of SSI/ Medicare days for FY 1985 derived from matching FY 1985 SSI eligibility files to Medicare FY 1985 PATBILL records.

In accomplishing the standardization, we did not take into account any payments to hospitals that qualify for disproportionate share payments based on the percentage of their revenue from State and local government sources for indigent care. This is because these hospitals must demonstrate on a hospital-by-hospital basis that they meet the criteria for a payment adjustment. We did not know at the time of the publication of the proposed rule how many or which hospitals would ultimately qualify under this provision. While it was our belief that the number of these hospitals would be small, and therefore would not have a significant effect on the standardized rates, we stated that we would monitor this situation closely, and, to the extent possible, present our data and analysis in the final rule. We stated that, if a larger number of hospitals than we expected do qualify, we would consider restandardizing the rates as a part of the final rule to take account of payments to these hospitals. However, currently. there are still no hospitals that have qualified for disproportionate share payments under this provision.

We also noted in the proposed rule that section 9306(a) of Pub. L. 99-509 amended section 1886(d)(5)(F)(v) of the Act to provide that a hospital that is located in a rural area and has 500 or more beds also serves a significantly disproportionate number of low-income patients for a cost reporting period if the hospital has a disproportionate patient percentage that equals or exceeds a percentage specified by the Secretary. We stated in the proposed rule that if standardization is necessary to take into account additional payments as a result of that rulemaking, we would do it as part of this final rule. This provision of the law was implemented through a final rule that was published in the Federal Register on June 25, 1987 (52 FR 23832). In the impact analysis prepared as a part of that rule, we estimated that only two hospitals would qualify as disproportionate share hospitals under the rule. We have now determined that only one hospital will qualify. Therefore, we do not believe that it is necessary to make any adjustment to the payment rates since the effect is negligible.

2. Grouping of Urban/Rural Averages Within Geographic Areas. Under section 1886(d)(2)(D) of the Act, the average standardized amounts must be determined for hospitals located in urban and rural areas of the nine census divisions and the nation. For FY 1988, the Federal rates will be comprised of 100 percent of the national rate (section 1886(d)(1)(A)(iii) of the Act). Section 1886(d)(5)(C)(ii) of the Act specifies that a sole community hospital's Federal rate is based on 100 percent of the regional rate.

In previous prospective payment final rules, Table 1 has contained 20 standardized amounts (ten urban amounts and ten rural amounts which are further divided into labor-related and nonlabor-ralated portions). However, this year we are splitting Table 1 into Tables 1a and 1b. (Table 1c applies to Puerto Rico, as discussed below.) Table 1a would contain the two national standardized amounts that are applicable to most hospitals. Table 1b would set forth the 18 regional standardized amounts applicable to sole community hosptials. The methodology for computing the national average standardized amounts is identical to the methodology for determining the regional amounts, except that the national urban and rural groups include hospitals from all urban and all rural geographic areas, respectively.

Currently, the average standardized amounts are based on hospital-weighted averages; that is, the average standardized amount is the average of the average standardized costs per discharge of all hospitals. As a result, each hospital, regardless of its number of discharges, has an equal impact on the average.

Section 9302(c) of Pub. L. 99–509 amended section 1886(d)(3)(A) of the Act to specify that, with respect to discharges occurring on or after October 1, 1987, urban and rural averages are to be computed on the basis of discharge-weighting rather than hospital-weighting. Under discharge-weighting, the standardized amounts are based on an average derived by dividing total costs by the number of discharges. Thus, a hospital with a high number of discharges has a correspondingly greater impact on the overall average.

Section 1886(d)(3)(A) of the Act also specifies that appropriate adjustments are to be made to ensure that average standardized amounts computed on the basis of discharge-weighting do not result in total payments that are greater or less than the total payments that would have been made had the average standardized amounts been computed on the basis of hospital-weighting; that is, this provision must be "budget neutral". (For a detailed discussion of budget neturality, see section IV of this addendum.)

The Executive Office of Management and Budget (EOMB) has not announced any revised listings of the Metropolitan Statistical Area (MSA) and New England County Metropolitan Area (NECMA) designations that are used in calculating the standardized amounts. Therefore, the designations of MSAs and NECMAs contained in the wage

index set forth in the proposed rule will remain the same for this final rule.

Comment: We received one comment concerning discharge-weighting. The commenter was concerned that using discharge-weighting in computing the DRG weighting factors is disadvantageous to hospitals with high case-mix index values. In particular, the commenter noted that the relative weights of some high-cost DRGs, such as those for burn cases, have been significantly reduced by the recalibration.

Response: We believe that the commenter has confused the revised computation of the average standardized amounts with the calculation of the DRG relative weights. Weighting the operating cost per case for each hospital in the prospective payment system data base by its volume of discharges has the effect of increasing the average standardized amounts because it gives greater weight to highvolume hospitals, which tend to be more expensive. The increased average standardized amounts are then uniformly adjusted to ensure budget neutrality of total payments relative to estimated payments that would have been made based on rates that were hospital-weighted. These adjustments, however, are limited to the average standardized amounts and do not enter into either the DRG classification or the

recalibration of the DRG weights.

Recalibration of the DRG relative weights, on the other hand, is carried out in accordance with the steps described in section II.B. of the preamble of this final rule. As described in that section, the relative weights are normalized. That is, the average case weight for the FY 1986 MEDPAR cases using the revised DRG definitions and recalibrated weighting factors is computed. Similarly, the average case weight for the FY 1986 discharges using the current DRG definitions and weighting factors is computed. The ratio of the latter average case weight to the former is a normalization factor, which is then applied uniformly to the relative weights. Application of this normalization factor ensures that the average case weight for the FY 1986 MEDPAR cases used is constant before and after reclassification and recalibration, thus ensuring that reclassification and recalibration neither increase nor decrease estimated Medicare outlays for the set of cases on which the weights are based.

The final relative weight for each DRG represents the ratio of the average resources used to treat cases in the DRG to the average resources used to treat all Medicare cases in all DRGs. Changes in

the weighting factor for each DRG reflect classification changes (if any were made) and the relationship between changes in the average standardized charges for cases in each DRG and changes in the average standardized charges of all other DRGs.

With respect to the commenter's particular concern with the weighting factors of the burn DRGs, our data indicate that the average standardized charges for these DRGs have declined from FY 1984 to FY 1986. We believe that it is this decline in charges, rather than the change to discharge-weighting, which accounts for the weighting factors for these DRGs.

3. Updating the Average Standardized Amounts. In accordance with section 1886(d)(3)(A) of the act as amended by section 9302(a)(2) of Pub. L. 99-509, we are proposing to update the urban and rural average standardized amounts using the applicable percentage increase specified in section 1886(b)(3)(B) of the Act, as amended by section 9302(a)(1) of Pub. L. 99-509. The percentage increase to be applied is mandated under that section of the law as the estimated increase in the hospital market basket percentage minus 2.0 percentage points. The percentage change in the market basket reflects the average change in the price of goods and services purchased by hospitals to furnish inpatient care.

In the September 3, 1986 final rule, we revised the hospital market basket by rebasing to reflect 1982, rather than 1977, cost data, expanding the number of market basket cost categories from 18 to 28, and modifying certain variables used as the price proxies for some of the cost categories. For a detailed discussion of this revision, see 51 FR 31461–31468.

When the proposed rule was published, the increase in the hospital market basket was estimated at 4.7 percent. Therefore, the proposed applicable percentage increase was 2.7 percent (market basket percentage increase minus 2.0 percentage points). Thus, we proposed that the standardized amounts and the hospital-specific rates (which for cost reporting periods beginning on or after October 1, 1987 apply only to sole community hospitals) be increased by 2.7 percent.

Although the update factor for FY 1988 is set by law, we are required by section 1886(e)(4) of the Act to recommend an appropriate update factor for FY 1988. Under section 1886(e)(5) of the act, we are required to publish both our proposed and final recommendations of an update factor. We published our proposed recommendation in the Federal Register

on June 11, 1987 (52 FR 22386). Our final recommendation is set forth in Appendix B of this final rule.

Comment: We received a number of comments concerning the proposed update factor of 2.7 percent (market basket percentage increase minus 2.0 percent). Many commenters supported this update factor while some believe that the rates should be increased by the full market basket increase.

Commenters were also concerned that we would implement the proposed recommended update of 0.75 percent set forth in the June 11, 1987 notice rather than the 2.7 percent set forth in the proposed rule.

Response: The percentage increase to be applied to the rates for FY 1988 is mandated by section 1886(d)(3)(A) of the Act as amended by section 9302(a)(2) of Pub. L. 99–509. This section of the law specifies the update for FY 1988 as the estimated increase in the hospital market basket minus 2.0 percentage points. The most recent forecast of the market basket increase for FY 1988 remains at 4.7 percent. Therefore, the applicable percentage increase is 2.7 percent.

Since the update for FY 1988 is set by law, we do not have the authority to apply a different percentage increase from that prescribed in the law. However, section 1886(e)(4) of the Act, as amended by section 9302(a)(2)(B) of Pub. L. 99-509, requires that the Secretary, taking into consideration the recommendations of ProPAC. recommend an appropriate update factor for FY 1988, which takes into account amounts necessary for the efficient and effective delivery of medically appropriate and necessary care of high quality. Accordingly, in the June 11, 1987 notice, we recommended an update of 0.75 percent for prospective payment hospitals and 1.9 percent for hospitals excluded from the prospective payment system. We note that these updates are only our recommendations and cannot be implemented without Congressional action. The comments we received in response to our proposed recommendation are addressed in Appendix B of this final rule.

Comment: One commenter expressed concern that the HCFA Hospital Occupational Index used in forecasting the hospital market basket increase does not reflect the unique labor market from which hospitals must recruit workers. This is because the labor component of the market basket is measured using a blend of hospital and nonhospital wage indicators with nonhospital wages accounting for nearly

three-fourths of the labor component. Previously (prior to FY 1987), the labor component of the market basket was based solely on hospital wage trends. This commenter recommended that HCFA revert to the exclusive use of hospital industry wage data for the labor component of the market basket and suggested the use of the Employment Cost Index (ECI) for hospitals, which was developed by the Bureau of Labor Statistics.

Response. As discussed above, in the September 3, 1986 final rule, we revised the hospital market basket by rebasing to reflect 1982, rather than 1977, cost data. As part of that rebasing, we modified the variables used as price proxies for the labor component of the hospital market basket. We developed the HCFA Hospital Occupational Index, which uses a combination of hospital and nonhospital wage proxies.

In its April 1, 1985 report to the Secretary, ProPAC expressed concern that use of the BLS average hourly earning index for hospitals used in the previous market basket did not distinguish between changes in inflation and changes in occupational mix. That is, rapid increases in average hourly wages could reflect changes in skill mix instead of in wage rates. ProPAC suggested that a combination of internal and external (hospital and nonhospital) proxies should be used to measure

changes in wages.

The issue of whether to use only an internal wage proxy (that is, one based exclusively on hospital wage and salary data) or a combination of internal and external wage proxies has been debated for some time. The market basket is intended to measure prices actually faced by the hospital industry. Thus, for labor, we wish to measure only changes in wage rates, not changes in the composition of the labor used by hospitals. In rebasing the market basket, we decided to use an external measure in addition to an internal measure because the external measure (the employment cost index) reflects changes in the price of wages only instead of changes in wage prices and labor mix, as reflected by the internal measure (Average Hourly Earnings of Hospital Workers).

We indicated in the September 3, 1986 final rule that once an employment cost index specific to hospital workers becomes available we would consider using it rather than the current blend of internal and external measures (54 FR 31465). As the commenter noted, the employment cost index for hospital worker categories has been recently developed by the Bureau of Labor Statistics. However, we believe its use

as a measure of hospital labor price changes is premature at this time because there are not yet a sufficient number of historical observations from which to base accurate forecasts.

4. Other Adjustments to the Average Standardized Amounts-a. Part B Costs. Section 1862(a)(14) of the Act prohibits payments for nonphysician services furnished to hospital inpatients unless the services are furnished either directly by the hospital, or by an entity under arrangements made by the hospital under which Medicare's payment to the hospital discharges the beneficiary's liability to pay for the services furnished.

In the September 3, 1985 final rule, we increased the average standardized amounts by 0.13 percent so that they represent costs previously billed under Part B (50 FR 35708). In the September 3, 1986 final rule, we stated that we were making no further adjustments for this factor in FY 1987, or in future Federal fiscal years, because the appropriate adjustment had been built into the FY

1986 base (51 FR 31521).

b. FICA Taxes. Section 1886(b)(6) of the Act requires that adjustments be made in the base period costs in recognition of the fact that certain hospitals were required to enter the Social Security system and begin paying FICA taxes as of January 1, 1984. In the September 3, 1985 final rule, we increased the average standardized amounts by 0.18 percent to account for additional costs of payroll taxes for hospitals entering the Social Security system (50 FR 35708). In the September 3, 1986 final rule we stated that we were making no further adjustments for this factor in FY 1987, or in future Federal fiscal years, because the appropriate adjustment has also been built into the FY 1986 base.

c. Nonphysician Anesthetist Costs. Section 1886(d)(5)(E) of the Act provides that hospital costs for the services of nonphysician anesthetists are paid in full as a reasonable cost pass-through. Under section 2312(c) of Pub. L. 98-369. this pass-through was made effective for cost reporting periods beginning on or after October 1, 1984, and before October 1, 1987. Section 9320(a) of Pub. L. 99-509 extended the period of applicability of this pass-through so that services will continue to be paid under reasonable cost for any cost reporting periods (or parts of cost reporting periods) ending before January 1, 1989 and struck subsection (E) effective on

In the September 3, 1985 final rule, we noted that to the extent an adjustment was warranted to reflect the removal of these costs from the prospective

payment rates for FY 1985, it was incorporated in the overall budget neutrality adjustment (50 FR 35708). Therefore, because this adjustment has already been built into the FY 1985 base from which the FY 1986, FY 1987, and proposed FY 1988 rates are derived, we did not propose to make further adjustments to the average standardized amounts for FY 1988.

d. Indirect Medical Education. Section 9104(b) of Pub. L. 99-272 added section 1886(d)(3)(C)(ii) to the Act to provide that, effective for discharges occurring on or after October 1, 1986, the average standardized amounts be further reduced, taking into consideration the effects of the standardization for indirect medical education costs as described in section II.A.1.c. of this addendum. Specifically, for each geographic area (regional and national, urban and rural), total payments including indirect medical education and disproportionate share hospital adjustments, based on payment rates standardized for an 8.1 percent curvilinear indirect medical education factor and for disproportionate share, shall be neither more nor less than the estimated total of payments, including indirect medical education adjustment payments that would have been made based on rates standardized for an 11.59 percent linear indirect medical education factor and paid out at 8.7 percent on a curvilinear basis. The adjustment is accomplished on a regional basis in order to reflect congressional intent that the necessary calculations will not redistribute payments among the regions. Through this adjustment, Congress is ensuring that total prospective payments, on a regional basis, taking into consideration the restandardization of rates for disproportionate share payments and for a revised indirect medical education payment factor of approximately 8.1 percent on a curvilinear basis, will equal payments that would have resulted with rates standardized for an 11.59 percent linear indirect medical education adjustment factor, and payments computed using an indirect medical education factor of 8.7 percent applied on a curvilinear basis. For discharges on or after October 1, 1989 (that is, after that part of the law requiring disproportionate share payments ceases to be in effect), the adjustment must be such as to ensure that the system savings resulting from the changes to the indirect medical education factor are preserved.

Therefore, under section 1886(d)(3)(C)(ii) of the Act, for FY 1988 we proposed to adjust the urban and

rural regional and national standardized amounts to account for indirect medical education payments. This adjustment is made in conjunction with the budget neutrality adjustments (see section IV of this addendum).

f. Outliers. Section 1886(d)(5)(A) of the Act requires that, in addition to the basic prospective payment rates, payments must be made for discharges involving day outliers and may be made for cost outliers. Section 1886(d)(3)(B) of the Act correspondingly requires that the standardized amounts be reduced by the proportion of estimated total DRG payments attributable to estimated outlier payments. Furthermore, section 1886(d)(5)(A)(iv) of the Act directs that outlier payments may not be less than five percent nor more than six percent of total payments projected to be made based on the prospective payment rates in any year. For FY 1987, we set the outlier thresholds so as to result in estimated outlier payments equal to five

percent of total payments.

Section 9302(b)(1) of Pub. L. 99-509 amended section 1886(d)(3)(B) of the Act to require that, effective with discharges occurring on or after October 1, 1986, each national and regional standardized amount be reduced for hospitals located in urban areas and for hospitals located in rural areas based on the estimated proportion of total DRG payments attributable to outlier payments for hospitals in urban areas and for hospitals in rural areas, respectively. Consequently, instead of the uniform five percent reduction factor applying equally to all the standardized amounts. there are now two separate reduction factors, one applicable to the urban national and regional standardized amounts and the other applicable to the rural national and regional standardized amounts. Rates for urban hospitals, which are projected to receive outlier payments in excess of five percent of total DRG payments, are reduced by that larger percentage (instead of by five percent). Rates for rural hospitals, which are projected to receive outlier payments of less than five percent of total DRG payments, are reduced by the lower percentage (instead of by five percent).

We proposed to continue to set the outlier thresholds so as to result in estimated outlier payments equal to five percent of total prospective payments. Therefore, for FY 1988, we proposed to set the day outlier threshold at the lesser of 23 days or 2.0 standard deviations and the cost outlier threshold at the greater of 2.0 times the prospective payment rate for the DRG or \$16,000.

The proposed outlier reduction factors for FY 1988 were as follows:

Outlier Reduction Factors

Urban: .94519. Rural: .97246.

As indicated in section V.C. of the preamble to this final rule, we have decided, based on comments received. not to adopt certain charges in outlier payment methodologies that we had proposed for FY 1988. These proposed changes affected the level of the proposed outlier thresholds. As a result of not implementing the proposed changes, as well as the incorporation of later charge data for FY 1986, we are setting the day outlier threshold at the lesser of 18 days or 2.0 standard deviations and the cost outlier threshold at the greater of 2.0 times the prospective payment rate for the DRG or \$14,000.

The final outlier reduction factors for FY 1988 are as follows:

Outlier Reduction Factors

Urban: .94441 Rural: .97485

In another document 1 we are revising the prospective payment system to incorporate capital costs. Under the provisions of that document, payments may be made for outliers as a part of the capital payment in the same way that have been made for inpatient operating costs since the implementation of the prospective payment system. This final rule on capital costs contains the methodology, as well as examples, of how we compute the capital component of the outlier payment. This system for incorporating capital will be effective with cost reporting periods beginning on or after October 1, 1987. Until a hospital becomes subject to the new capital system, payment for outliers will be made using only the current methodology. (See the examples in the September 3, 1986 final rule (51 FR

B. Adjustments for Area Wage Levels and Cost-of-Living

This section contains an explanation of the application of two types of adjustments to the adjusted standardized amounts that will be made by the intermediaries in determining the prospective payment rates as described in section D below. For discussion purposes, it is necessary to present the adjusted standardized amounts divided into labor and nonlabor portions. Tables la and lb contain the actual labor-related and nonlabor-related shares that

will be used to calculate the prospective payment rates for hospitals located in the 50 States and the District of Columbia.

1. Adjustment for Area Wage Levels. Section 1886(d)(2)(H) of the Act requires that an adjustment be made to the laborrelated portion of the prospective payment rates to account for area differences in hospital wage levels. This adjustment is made by the intermediaries by multiplying the laborrelated portion of the adjusted standardized amounts by the appropriate wage index for the area in which the hospital is located. In section III of the preamble to this final rule, we discuss certain revisions we are making to the wage index. This index is set forth in Tables 4a and 4b of this addendum.

2. Adjustment for Cost of Living in Alaska and Hawaii. Section 1886(d)(5)(C)(iv) of the Act authorizes an adjustment to take into account the unique circumstances of hospitals in Alaska and Hawaii. Higher laborrelated costs for these two States were included in the adjustment for area wages above. For FY 1988, the adjustment necessary for nonlaborrelated costs for hospitals in Alaska and Hawaii will be made by the intermediaries by multiplying the nonlabor portion of the standardized amounts by the appropriate adjustment factor contained in the table below.

Table of cost-of-Living Adjustment Factors, Alaska and Hawaii Hospitals

Alaska-All areas	1.25
Hawaii:	
Oahu	1.225
Kauai	
Maui	
Molokai	1.20
Lanai	1.20
Hawaii	1.15

(The above factors are based on data obtained from the U.S. Office of Personnel Management.)

C. DRG Weighting Factors

As discussed in section II of the preamble to this final rule, we have developed a classification system for all hospital discharges, sorting them into DRGs, and have developed weighting factors for each DRG that are intended to reflect the relative average resource consumption associated with each DRG.

Table 5 of section VI of this addendum contains the weighting factors that we will use for discharges occurring in FY 1988. These factors have been recalibrated as explained in section II of the preamble.

¹Editorial note.—The capital costs final rule appears in Part IV of this issue.

D. Calculation of Prospective Payment Rates for FY 1988

General Formula for Calculation of Prospective Payment Rates for Cost Reporting Periods Beginning on or after October 1, 1987 and Before October 1, 1988

Prospective Payment Rate for all hospitals except sole community hospitals equals Federal Portion (100 percent national rate)

Prospective Payment Rate for Sole Community Hospitals equals 75 percent of the hospital-specific portion plus 25 percent of the Federal portion (100 percent regional

1. Federal Portion. For cost reporting periods beginning on or after October 1, 1987 and before October 1, 1988, except for sole community hospitals, 100 percent of the hospitals rate is the hospital's Federal rate. Beginning with discharges occurring on or after October 1, 1987, the Federal rate is comprised of 100 percent of the Federal national rate except for sole community hospitals, whose 25 percent Federal portion is based on the Federal regional rate. The Federal rates are determined as follows:

Step 1-Select the appropriate regional or national adjusted standardized amount considering the type of hospital and urban or rural designation of the hospital (see Tables 1a and 1b, section VI of this addendum).

Step 2-Multiply the labor-related portion of the standardized amount by the appropriate wage index.

Step 3-For hospitals in Alaska and Hawaii, multiply the nonlabor-related portion of the standardized amount by the appropriate cost-of-living adjustment factor.

Step 4-Sum the amount from step 2 and the nonlabor portion of the standardized amount (adjusted if appropriate under step 3).

Step 5-Multiply the final amount from step 4 by the weighting factor corresponding to the appropriate DRG weight (see Table 5, section VI of this addendum).

2. Hospital-Specific Portion (Applicable only to Sole Community Hospitals). The hospital-specific portion of the prospective payment rate is based on a hospital's historical cost experience. For the first cost reporting period under prospective payment, a hospital-specific rate was calculated for each hospital, derived generally from the following formula:

Base year costs per discharge divided by 1981 case-mix index times updating factor equals Hospital-specific rate

For sole community hospitals, the hospital-specific portion equals 75 percent of the hospital-specific rate for all cost reporting periods beginning on or after October 1, 1983. For each

subsequent cost reporting period, the hospital-specific portion is derived as

Hospital-Specific Rate times Updating Factor times Blending Percentage (75 percent) times DRG Weight.

For a more detailed discussion of the hospital-specific portion, we refer the reader to the September 1, 1983 interim final rule (48 FR 39772).

a. Updating the Hospital-Specific Rates for FY 1988 Cost Reporting Periods. We are increasing the hospitalspecific rates by 2.7 percent (market basket percentage increase minus two percentage points) for cost reporting periods beginning on or after October 1, 1987. As required by sections 1886(b)(3)(A) and (B) of the Act (as amended by section 9302 of Pub. L. 99-509), this is the same percentage increase (2.7 percent) by which we are changing the Federal rates for FY 1988.

b. Calculation of Hospital-Specific Portion. For sole community hospital cost reporting periods beginning on or after October 1, 1987, the hospitalspecific portion of a hospital's payment for a given discharge is calculated by-

Step 1-Multiplying the hospital's hospital-specific rate by the applicable update factor (1.027);

Step 2-Multiplying the result in Step

1 by 75 percent; and

Step 3-Multiplying the amount resulting from Step 2 by the specific DRG weighting factor applicable to the discharge. The result is the hospitalspecific portion of the FY 1988 prospective payment for a given discharge for a sole community hospital.

III. Prospective Payment Rates for Hospitals Located in Puerto Rico

This section contains an explanation of how we derive the adjusted standardized payment amounts applicable for FY 1988 for hospitals located in Puerto Rico. The methodology for arriving at the appropriate rate structure is essentially prescribed by section 1886(d)(9) of the Act and is set forth in regulations in §§ 412.207 through

A. Calculation of Adjusted Standardized Amounts

The Puerto Rico adjusted standardized amounts, which are set forth in Table 1c, are computed as described below.

 Target Amounts. Section 1886(d)(9)(B)(i) of the Act requires that we determine the Medicare target amount (as defined in section 1886(b)(3)(A) of the Act) for each hospital for its cost reporting period beginning in FY 1987. For purposes of

computing the Puerto Rico standardized amounts, we will not consider revisions to the target amounts subsequent to HCFA's development of those amounts.

2. Updating for FY 1988. Section 1886(d)(9)(B)(i) of the Act also requires that each target amount be updated to the midpoint of FY 1988 (April 1, 1988) by prorating the applicable percentage increase for FY 1988 as defined in section 1886(b)(3)(B) of the Act. That section of the Act specifies that the applicable percentage increase for FY 1988 is the increase in the market basket percentage minus 2.0 percentage points, that is, 2.7 percent.

3. Standardization of the Target Amount. Section 1886(d)(9)(B)(ii) of the Act requires that the updated target amount for each hospital be standardized for several variables. Standardization means the removal of the effects of certain sources of variation in cost among hospitals. These include case mix, differences in area wage levels, payments for hospitals that serve a disproportionate share of lowincome patients, and indirect medical education costs.

a. Adjustments for Variations in Hospital Wage Levels. Section 1886(d)(9)(B)(ii)(II) of the Act requires that the updated target amount be standardized by adjusting for variations among hospitals by area in the average area hospital wage level. Therefore, the target amount is divided into labor and nonlabor portions, based on the labor and nonlabor components of the hospital market basket. The laborrelated portion is then divided by the appropriate wage index for the geographic area in which the hospital is located to remove the effects of local wage differences from hospital target amounts.

As discussed in section III of the preamble, we are updating the HCFA wage index using 1984 data and making a change in the methodology for computing the national average hourly wage, which serves as the basis for indexing the area wage levels. In addition, as discussed in section IV of the preamble, we are adding wage index values for areas in Puerto Rico to the wage index. The wage index is set forth in Tables 4a and 4b.

b. Variations in Case Mix Among Hospitals. Section 1886(d)(9)(B)(ii)(III) of the Act requires that the updated target amounts be standardized to adjust for variations in case mix among hospitals. The methodology used for determining the appropriate adjustment factor (that is, the case-mix index) is explained in the September 1, 1983 interim final rule (48 FR 39768-39771). A case-mix index

has been calculated for each hospital in Puerto Rico based on 1984 data.

Standardization, necessary to neutralize inpatient operating costs for the effects of variations in case mix, is accomplished by dividing the hospital's target amount per Medicare discharge by that hospital's case-mix index.

c. Indirect Medical Education Costs.
Section 1886(d)[9)(B)(ii)(I) of the Act
requires that the updated target amounts
be standardized for indirect medical
education costs. Section 1886(d)[9)(D)(ii)
of the Act provides that prospective
payment hospitals in Puerto Rico receive
an additional payment for the indirect
costs of medical education as specified
in section 1886(d)(5)(B) of the Act. Under
section 1886(d)(5)(B) of the Act, the
indirect medical education cost payment
is based on an education adjustment

factor, which is approximately 8.1 percent for discharges occurring on or after May 1, 1986 and before October 1, 1989. For discharges occurring on or after October 1, 1989, the adjustment factor is equal to approximately 8.7 percent. These factors are approximations because the adjustment factor is calculated on a curvilinear or variable basis. An adjustment made on a curvilinear basis reflects a nonlinear cost relationship, that is, each absolute increment in a hospital's ratio of interns and residents to beds does not result in an equal proportional increase in costs. Therefore, the adjustment factors are only approximately 8.1 percent and 8.7

For discharges occurring on or after May 1, 1986 and before October 1, 1989, the indirect medical education factor is calculated using the following formula:

For discharges occurring on or after October 1, 1989, the indirect medical education factor equals the following:

Therefore, after adjusting each hospital's updated target amount for differences in area wage levels and case mix, we divided each teaching hospital's target amount by 1.0 plus the individual hospital's indirect medical education adjustment factor as computed using the formula described above, which section 1886(d)(5)(B)(ii)(I) of the Act requires be used for discharges on or after May 1, 1986 and before October 1, 1989.

d. Costs for Hospitals That Serve a Disproportionate Share of Low-Income Patients. Section 1886(d)(9)(B)(ii)(IV) of the Act provides that the updated target amounts be standardized for the estimated additional payments made to hospitals that serve a disproportionate share of low-income patients. That is, the law requires us to remove the effects of the payments made to disproportionate share hospitals from

the costs used to establish the standardized amounts.

Therefore, we are adjusting each disproportionate share hospital's updated target amount by adding 1.0 to the applicable disproportionate share payment factor, and dividing the hospital's updated target amount by that number. In this way, we remove the effect of payment adjustments for disproportionate share hospitals from the standardized amounts as required under section 1886(d)(9)(B)(ii)(IV) of the Act.

In determining the disproportionate share adjustment factors for purposes of standardizing the updated target amounts, we will use available data on the percentage of Medicaid days from FY 1984 Medicare cost reports and the percentage of SSI/Medicare days for FY 1985 derived from matching FY 1985 SSI

eligibility files to Medicare FY 1985 PATBILL records.

In accomplishing this standardization, we have not taken into account any payments to hospitals that qualify for disproportionate share payments based on the percentage of their revenue from State and local government sources for indigent care. This is because these hospitals must demonstrate on a hospital-by-hospital basis that they meet the criteria for a payment adjustment.

4. Grouping of Urban/Rural Averages Within Geographic Areas. Under section 1886(d)(9)(B)(iii) of the Act, the average standardized amount per discharge must be determined for hospitals located in urban and rural areas in Puerto Rico. That section of the Act also specifies that the urban and rural average standardized amounts for Puerto Rico hospitals are based on discharge-weighted averages just as section 1886(d)(3)(a) of the Act specifies this methodology for the average standardized amounts that are applicable to other prospective payment hospitals. This methodology is discussed in detail in section II.A.2. of this addendum. The average standardized amounts for hospitals located in Puerto Rico are set forth in Table 1c.

5. Other Adjustments to the Average Standardized Amounts. The average standardized amounts, calculated as described above, are further adjusted as explained below. Note that there are no adjustments for Part B costs or FICA taxes for hospitals located in Puerto Rico as there are for prospective payment hospitals located outside of Puerto Rico. This is because adjustments to account for these costs have already been made to the target amounts on which the average standardized amounts are based.

a. Nonphysician Anesthetist Costs. Section 1886(d)(9)(D)(iv) of the Act specifies that the provisions of section 1886(d)(5)(E) of the Act apply to hospitals located in Puerto Rico. Section 1886(d)(5)(E) of the Act provides that hospital costs for the services of nonphysician anesthetists are paid in full as a reasonable cost pass-through. Under section 2321(c) of Pub. L. 98-369, this pass-through was made effective for cost reporting periods beginning on or after October 1, 1984, and before October 1, 1987. Section 9320(a) of Pub. L. 99-509 extended the period of applicability of this pass-through so that services will continue to be paid under reasonable cost for any cost reporting periods (or parts of cost reporting periods) ending before January 1, 1989 and struck subsection (E) effective on that date.

We considered the effect of the passthrough provision on the average adjusted standardized amounts as part of the budget neutrality analysis (see discussion in section IV of this

addendum).

b. Outliers. Section 1886(d)(5)(A)(iv) of the Act, made applicable to Puerto Rico by section 1886(d)(9)(D)(i) of the Act, directs that outlier payments may not be less than five percent nor more than six percent of total payments projected to be made to prospective payment hospitals based on the payment rates in any year. Since Puerto Rico hospitals will be subject to the prospective payment system beginning October 1, 1987, bills from those hospitals have been used in setting the proposed outlier thresholds (set forth above in section II.A.4.f. of the addendum) so that overall system-wide outlier payments are estimated to be five percent of total prospective payments as required by law.

Section 1886(d)(3)(B) of the Act requires that separate urban and rural outlier offsets to the standardized amounts be developed. As initially implemented October 1, 1986, these offsets apply on a national basis to urban and rural hospitals. We proposed to set the same outlier offsets for the Puerto Rico prospective payment standardized amounts as we have for hospitals located outside Puerto Rico. The proposed outlier reduction factors

were as follows:

Urban	Rural
.94519	.97246

The final reduction factors for FY 1988 are as follows:

Urban	Rural
.94441	.97485

B. Calculation of National Standardized Amount for Puerto Rico

The national standardized payment amount applicable to hospitals in Puerto Rico consists of the discharge-weighted average of the national rural standardized amount and the national urban standardized amount (as set forth in Table 1a of this addendum). The national average standardized amount for Puerto Rico is set forth in Table 1c.

C. Adjustments for Area Wage Levels

Section 1886(d)(9)(B)(vi) of the Act requires that an adjustment be made to

the labor-related portion of the Puerto Rico prospective payment rates to account for area differences in hospital wage levels. This adjustment is made by the intermediaries by multiplying the labor-related portion of the adjusted standardized amounts by the appropriate wage index for the area in which the hospital is located. (Table 1c sets forth the labor-related and nonlabor-related shares for both the Puerto Rico and the national standardized amounts that would be used to calculate the prospective payment rates for hospitals located in Puerto Rico.) The wage index is set forth in Tables 4a and 4b of this addendum.

D. DRG Weighting Factors

As discussed in section II of the preamble to this final rule, we have developed a classification system for all hospital discharges, sorting them into DRGs, and have developed weighting factors for each DRG that are intended to reflect the relative resource consumption associated with each DRG.

Table 5 of section VI of this addendum contains the weighting factors that we will use for discharges occurring in FY 1988. These factors have been recalibrated as explained in section II of the preamble.

E. General Formula for Calculation of Prospective Payment Rates for Hospitals Located in Puerto Rico Beginning On or After October 1, 1987 and Before October 1, 1988

Prospective Payment Rate for Puerto Rico hospitals = 75 percent of the Puerto Rico Rate + 25 percent of the National Rate.

 Puerto Rico Rate. The Puerto Rico prospective payment rate is determined as follows:

Step 1—Select the appropriate adjusted average standardized amount considering the urban and rural designation of the hospital (see Table 1c, section VI of the addendum).

Step 2—Multiply the labor-related portion of the standardized amount by the appropriate wage index.

Step 3—Sum the amount from step 2 and the nonlabor portion of the standardized amount.

Step 4—Multiply the amount from step 3 by the weighting factor corresponding to the appropriate DRG weight (see Table 5, section VI of the addendum).

2. National Rate. The national prospective payment rate is determined as follows:

Step 1—Multiply the labor-related portion of the national average

standardized amount (see Table 1c, section VI of the addendum) by the appropriate wage index.

Step 2—Sum the amount from step 1 and the nonlabor portion of the national average standardized amount.

Step 3—Multiply the amount from step 2 by the weighting factor corresponding to the appropriate DRG weight (see Table 5, section VI of the addendum).

IV. Budget Neutrality

The law requires that a number of adjustments be made to the average standardized amounts in order to achieve the payment levels anticipated by Congress in its revisions to section 1886 of the Act. In order to incorporate these adjustments, which are discussed in more detail below as well as in previous prospective payment rules, we used an iterative simulation process.

Using the most current data available (that is, bills for FY 1986 discharges from hospitals currently subject to the prospective payment system received in HCFA through June 1987 (approximately 9.7 million discharges)), we ran a baseline simulation using the PRICER program to price each case.

Estimated payments were calculated using FY 1988 standardized amounts computed on the same basis as those published in the September 3, 1986 final rule (51 FR 31530), except that these rates were—

 Updated by 1.15 percent for FY 1987 (rather than by .5 percent as announced in the September 3, 1986 final rule) and further updated by 2.7 percent for FY 1988 as prescribed by section 1886(b)[3)(B)(i)(II) of the Act;

 Adjusted to reflect the restandardization of the wage index resulting from revising the methodology for computing the national average hourly wage; and

 Adjusted to take into account the additional payments to rural referral centers as required by section 1886(d)[5](C)(i) of the Act.

The September 3, 1986 rates already included adjustments required by various provisions of Pub. L. 99–272, such as restandardization for indirect medical education payments, standardization for payments to hospitals serving a disproportionate share of low-income patients, and the adjustment for the indirect medical education payment equality factor (see 51 FR 31498–31529).

From this simulation, we calculated the ratio of total outlier payments to total payments (including outliers). We computed separate outlier payment ratios for hospitals in urban areas and hospitals in rural areas.

In addition, we calculated the total operating payments under the prospective payment system that we estimate would have occurred in FY 1988 using standardized amounts that were hospital-weighted and reduced uniformly for outliers by five percent. This amount served as the aggregate prospective payment target that had to be maintained after the urban and rural standardized amounts were dischargeweighted and differentially adjusted for urban and rural outlier ratios, respectively.

The next step was to discharge-weight the standardized amounts and to remove the effect of the five percent outlier adjustment from the FY 1988 standardized amounts and replace it with the initial outlier ratio estimates for urban hospitals and rural hospitals as computed in the initial price simulation. However, these initial outlier ratios require further refinements since they were computed on the basis of standardized payment amounts uniformly reduced by five percent. Therefore, further simulations were conducted to refine the outlier payment ratios used in computing the standardized amounts and to ensure that the total payment constraint was

These revised rates were also used to rerun the price simulation not only to refine the outlier payment ratios used to offset the standardized amounts but also to determine if aggregate payments based on these discharge-weighted, differentially adjusted rates equal the target payment amount computed in the baseline price simulation.

The entire simulation process was repeated until the outlier ratios and budget neutrality factor computed in the simulation and used to adjust the standardized rates resulted in total aggregate payments equal to the baseline target amount that represents our estimate of total prospective payment system payments for FY 1988 that would have been incurred had these provisions not been implemented.

The outlier adjustment and budget neutrality factors are as follows:

Outlier

Urban	Rural
.94441	.97485

Budget Neutrality Factor

97449

Section 1886(e)(1)(C) of the Act requires that the incorporation of hospitals in Puerto Rico into the prospective payment system in FY 1988 be accomplished in a budget-neutral fashion; that is, the aggregate payment to prospective payment hospitals including those located in Puerto Rico must be neither greater nor less than the payment amount that would have been made to those hospitals had section 9304 of Pub. L. 99-509, which added Puerto Rico hospitals to the prospective payment system, not been enacted. Accordingly, we analyzed what the total payment for FY 1988 would be if all prospective payment hospitals, including hospitals located in Puerto Rico, are paid under the prospective payment system and what the total payment for FY 1988 would be for these hospitals if the hospitals located in Puerto Rico are paid as if they are still subject to the rate-of-increase limits and all other hospitals receive their payment under the prospective payment system. The difference in payment amounts is considerably less than 0.1 percent, and, consequently, the budget neutrality adjustment for incorporating hospitals in Puerto Rico into the prospective payment system is negligible. Therefore, we believe that it is unnecessary to adjust the average standardized amounts to achieve budget neutrality.

V. Target Rate Percentages for Hospitals and Hospital Units Excluded From the Prospective Payment System

A. Background

The inpatient operating costs of hospitals and hospital units excluded from the prospective payment system are subject to rate-of-increase limits established under the authority of section 1886(b) of the Act, which is implemented in § 413.40 of the regulations. Under these limits, an annual target amount (stated as inpatient operating cost per discharge) is set for each hospital, based on the hospital's own cost experience. This target amount is applied as a ceiling on the allowable costs per discharge for the hospital's next cost reporting period.

A hospital that has inpatient operating costs per discharge in excess of its target amount will be paid no more than that amount. However, a hospital has inpatient operating costs less than its target amount will be paid its costs plus the lower of (1) 50 percent of the difference between the inpatient operating cost per dishcarge and the target amount, or (2) five percent of the target amount.

Each hospital's target amount is adjusted annually before the beginning of its cost reporting period, by an applicable target rate percentage for the 12-month period, prorated based on calendar year target rate percentages. For cost reporting periods beginning in FY 1983 and FY 1984, the applicable target rate percentage was the estimated hospital market basket increase factor plus one percentage point. For cost reporting periods beginning in FY 1985, the applicable target rate percentage was the estimated hospital market basket increase factor plus one-quarter of one percentage point. Under section 9101(e)(3) of Pub. L. 99-272, the applicable target rate percentage increase for cost reporting periods beginning on or after October 1, 1985 through September 30, 1986 is 5/24 of one percent. Section 9101 of Pub. L. 99-272 provides that for purposes of updating the target rate for FY 1987, the FY 1986 increase will be deemed to have been one-half of one percent. For cost reporting periods beginning in FY 1987. section 9302(a) of Pub. L. 99-509 provided that the applicable percentage increase was 1.15 percent.

B. Target Amounts for Cost Reporting Periods Beginning in FY 1988

For cost reporting periods beginning in FY 1988, under section 1886(b)(3)(i)(II) of the Act, as amended by section 9302(a) of Pub. L. 99-509, the applicable percentage increase is the market basket percentage increase minus 2.0 percentage points. Therefore, we proposed to increase each hospital's previous year's target amount by 2.7 percent. Thus, the same percentage increase applies to the target rate amounts for hospitals and units excluded from the prospective payment system as applies to the prospective payment rates for hospitals subject to that system. Since the most recent estimated increase in the market basket remains at 4.7 percent, each hospital's previous year's target amount will be increased by 2.7 percent for its cost reporting period beginning on or after October 1, 1987.

VI. Tables

This section contains the tables referred to throughout the preamble to this proposed rule and in this addendum. For purposes of this proposed rule, and to avoid confusion, we have retained the designations of Tables 1 through 5 that were first used in the September 1, 1983 initial prospective payment final rule (48 FR

39844). Tables 1a, 1b, 1c, 3c, 4a, 4b, 5, and 7 are presented below. The tables are as follows:

Table 1a—National Adjusted Standardized Amounts, Labor/Nonlabor Table 1b—Regional Adjusted Standardized Amounts, Labor/Nonlabor Table 1c—Adjusted Standardized Amounts

for Puerto Rico, Labor/Nonlabor

Table 3c—Hospital Case-Mix Indexes for Discharges Occurring in FY 1986

Table 4a—Wage Index for Urban Areas Table 4b—Wage Index for Rural Areas Table 5— Diagnosis-Related Groups, Table 7a—Length-of-Stay Percentiles Using FY 1987 DRG Classification

Table 7b—Length-of-Stay Percentiles Using FY 1988 DRG Classification

TABLE 1A.—NATIONAL ADJUSTED STAND-ARDIZED AMOUNTS, LABOR/NONLABOR

Urban		Rural		
Labor-related Nonla- bor- related		Labor- related	Nonla- bor- related	
2337.09	828.12	2123.20	587.97	

TABLE 1b.—REGIONAL ADJUSTED STANDARDIZED AMOUNTS, LABOR/NONLABOR1

	Urban		Rural	
	Labor- related	Nonlabor- related	Labor- related	Nonlabor- related
1. New England (CT, ME, MA, NH, RI, VT) 2. Middle Atlantic (PA, NJ, NY) 3. South Atlantic (DE, DC, FL, GA, MD, NC, SC, VA, WV) 4. East North Central (IL, IN, MI, OH, WI) 5. East South Central (AL, KY, MS, TN) 6. West North Central (IA, KS, MN, MO, NB, ND, SD) 7. West South Central (AR, LA, OK, TX) 8. Mountain (AZ, CO, ID, MT, NV, NM, UT, WY) 9. Pacific (AK, CA, HI, OR, WA)	2442.27 2214.47 2348.04 2477.09 2255.07 2346.62 2356.52 2248.43 2198.90	859.86 824.29 753.85 892.15 684.66 812.11 753.37 801.47 919.67	2350.23 2253.79 2155.54 2180.78 2136.28 2072.68 1990.81 2022.89 1958.41	696.88 657.49 572.20 634.8 533.4 569.39 524.17 606.48

Applicable Only to Sole Community Hospitals.

TABLE 1c.—ADJUSTED STANDARDIZED AMOUNTS FOR PUERTO RICO, LABOR/NONLABOR

Urban		Rural	
Labor- related	Nonlabor- related	Labor- related	Nonlabor- related
2046.38	367.93	1366.46	260.43

	Labor- related	Nonlabor- related
National	2285.09	769.74

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	CASE MIX 01.1676	-	SE M	2002	CASE M 01.413	ROVI 3006	CASE MIX 01.0132
010059	00.9123	010120	00.9295	020027	00.9123	030068	01.0476
	6	010122	.92	300	1.465	030071	000 3002
	892	N	.122	030003	1.176	030072	60
	323	010124	.068	3000	0.915	3007	15
	01.0479	010	26	300	1032	030074	34
	00-9266	1017	118	030008	276	030076	00.8571
	01.1212	010128	000	3000	1.232	030077	888
010069	-	N	.066	3001	.288	10	12
	01.1845	013		030011	.181		
010072	01.0790	010131	.098	030012	.072	030080	162
010073	9866 00	3	.877	030013	01-1225	030081	882
010074	95	3	.939	410080	01-1625	030082	3
010075	0	m	1.017	030016	01.1001	3008	610
010078	01-1523	m	161.	030017	9091-10	030084	
010019	01.0360	MY.	00-9290	030018	01.2159	030085	
010080	0	60	.412	030019	01.1058	030086	-
0	01.3651	010142	168	3002	272	030087	01.1226
	00.9939	~	01.0722	300	.262	030088	.187
-	01.1697		01.0839		01.2419	030089	01-1444
010085	01-1108		01.1108	030024	01.4398	030091	00.8714
010086	00.9243	•	01.0213		01.1554	040001	01.0491
010087	01.2418	010148	00.9548	30027	00.9728	040005	01.0551
010089	01.0389	010149	.131		01.3014	040003	0
0100010	01.0076	010150	5786.00	030033	01.0877	400040	01.2177
010000	01-2409	010153	0	030035	01-1677	040000	01.0535
010094	000	020001	200	250050	121	04007	
010095	076	020002	20	3003	01.5538	040008	4 54
010006	68	020004	01.0602	300	327	040009	01-0061
010097	- CP	020005	01.0110	030040	770	100	01.1191
010098	01.0357	020006	01.1160	030041	.902		000800
010099	01.0589	020007	01.0345	030043		040013	6600-10
001010	01-1782	020008	01.0635	4	966.	040014	34
101010		050009	00.8769	3004	1.048		54
010102	9778	020010	6.0	4	.913		00
010103	3210	020011	606.	030049	1.010		01-1663
010104	.3695	020012		030051	1.044		01-1356
010108	6650	020013	.876	030054	0.952	040019	01.0532
01010	1190	020014	.975		1.051		01.3028
011010	9019	020016		030057	1.231		01-1224
010111	9433	020017	960.	030059	.083	040022	01.1747
010112	036	2001	\$66°	0	1.136		5
010113	01.3286	2001	466.0	190060	.280	040025	956
911010	.145	005	.882	030062	01.0973	040026	01.2105
010115	946.	2002	.901	030063	1.13	040027	1.131
010117	01.0086	020024	2600-10	030064		040028	•07
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NOTE: CASE MIX INDEXES DO NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS. : CASE MIX INDEXES INCLUDE CASES RECEIVED IN HCFA CENTRAL OFFICE THROUGH JUNE., 1987.

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PROVIDER	CASE MIX	PROVIDER	- 111	~	ISE M	PROVIDER	S	~	W
30	. 9599	40100	Min	50046	1114	4	1.257	5016	
4002	0 0	401	L C	400	404	-	1 18	5016	
000	0	100	100	100	100	200	200	STOR STORY	
4003	6	10+		4000	+ PT -	100	10273	0100	
4003	6.	401	0	2005	0,982	_	1079	2016	7
4003	06.	401	0	5005	1 .130	100	1.08	5017	7
4003	-	401	report	5005	1.202		1.18	5017	-
003	.048	401	W.	5005	1.172	100	1 . 32	5017	-
04	00	401	0	050055	1.148	050113		501	NE
400	.023	103	gmd	5005	1.136		1.32	5017	-
4004	116	103		5005	1 - 226		1.25	5017	7
400	0	401	gar.	5005	1266	1000	1.31	5017	-
400	10	407	4 (2008	1.287		1 15	5018	
100		100	3 0	400	216		112	2018	4
100	20	104	7 6	0000	01701	4	77 00	070	4 5
400	5	TOS		0000	76701	- 0	04.00	0100	-
400	0	401		2000	1 . 542		1000	0100	-
400	-	105	CO.	2006	1.205		1.25	5018	
000	0	401	000	2006	10135	A.1	1.23	5018	0
000	6.	500	ped	2006	1.079	A.I	1.17	5018	
000	6.	500	-proft	5006	40401	A.	1.24	5018	
005	6.	500	21	5007	10175	P-4	1019	5019	0
000	5	500	100	5007	1.215	8.1	1.26	5019	
200	0	500	-	5007	1-206	- 0	1.46	5019	2
100	0	N C	4.0	1003	1 - 1 2 2	1 00	1.25	5019	9
000		200	O P	000	000	N A	200	20108	
000		200	mg /5	000	00000	A 4	7707	1000	9 5
900	70	200	0 ,	2000	60701	0 0	1013	2000	6
900	6	200	posit	2001	62401	W /	1015	2000	
900	60	200	DA I	2001	10412	200	1037	5019	9
900	6.	200	0	2001	1.219	MPS 4	1001	6109	0
900	00	200	R.	5007	10105	AC .	1017	2019	
900	0	500	0	2008	10179	ACS I	1045	5020	
100	60	500	C	5008	10341	APN.	1.20	2020	
700	-	500	6.71	5008	1.233	-4	1015	5020	
700	0	500	4.4	5008	1.337	4.50	1.25	5020	
100	0	500	-	5008	1.282	-304	1.22	5020	
100	0	500	V	5008	1.243	-94	1036	5020	0
007	60	500	N	5008	10101	-30	1.23	2050	
100	6.	500	N	5008	1.195	-94	0.95	5021	9
007	1.	500	10	2009	1.175	200	1.10	5021	
008	0	500	Desired.	5009	10149	-50	1.15	5021	9
800	6	500	0	2009	1.102	10	-	5021	
008	0	200	62.9	2009	10417	10	1.240	5021	
008	0.0	500	-	5005	1.089	10	1.238	5021	
008	6.	500	5	5009	1.169	LO.	1.373	5021	0
008	6 .	500	10	2009	1.245	10	1.121	5021	
008	10	500	E-1	5003	1.073	10	1.109	5022	
600	6.	500	0	SO.	1.294	10	6101	5022	
600	60	200		2010	1.700	10	1.432	5022	
600	6.	500	-	2010	1.253	10	1.158	5022	
040095	5	050043	01.4579	050102	01.2547	050161	01.2116	050225	01-1619
6004	-	200	pared.	2010	1.330	991050	1.197	5022	O PE
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NOTE: CASE MIX INDEXES DO NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS. : CASE MIX INDEXES INCLUDE CASES RECEIVED IN HOFA CENTRAL OFFICE THROUGH JUNE., 1987.

R CASE MIX.		01-1641	01.0850	01.2380		01.2423	00	01.1988	159	01.3996	00.9934	1681 010	0101377	00-8488	00.9484	715	01-4717	01.2518	01.2565	01.2439	010-104	01-1374	01.0943	01-1112	01-1377	01.0616	01.2353	01-1549	01-1513	01.3933	01.3886	00.9810	01-1432	01-2716	01.0801	01-1588	0102151	01.5186	0100010	01.2599	01.1059	01.1798	01.21.19
W	050526	25	25	5053	050531	53	050535	5053	23	050541	050542	4	050544	2 4	050547	050548	54	050520	050551	050552	05050	05050	050561	050562	050564	050566	050567	050568	050569	050570	050573	050575	050576	050577	050578	050580	050581	050583	505	505	200	202	050588
CASE MIX	01.5513	01.1668	1686-00	01-1366	00.600		00.9823	01-1386	9166-00	01.0318	01.6861	01.3332	01-3062	00.7997	01.2536	01.5284	01.2273	01.1854	00.9643	01.1195	01.02142	01.1175	01.0795	01.0999	01.2004	01-1575	01.3518	01.3180	01.1645	01.2139	01-1947	01-1041	01.0795	01.5733	00.8872	01.1692	9 6		01-1307	.162	-	16701	01.14.38
PROVIDER	050441	050442	050443	050444	050446	050447	050448	050449	050450	050451	050454	050455	0504000	050458	050459	050464	050467	050468	050469	050470	050473	050476	050477	050478	050481	050482	050485	050486	050487	050488	050491	050492	050494	050496	050497	050502	050503	050506	050510	050512	050515	050516	12050
CASE MIX	01.1167	.104	01-1005	.284	01-1604	.053	00.9975	01.5254	00.9580	01.2102	01.2712	01.2173	00.0155	01-2010	01.2168	00.8824	01.2771	01.3764	01.1117	01.4053	01-2121	01.0703	01.0060	01-1600	00.9692	01-1865	01-1173	00.7932	01-1499	01.0740	01.2005	01.2083	01.0536	01.3137	01-1937	00.9489	00.9020	01.1098	.296	19	0123	01.1099	01.00275
80	050371	050372	050373	050376	37	050378	2	3 3	050381	38	050383	050363	050387	050390	050391	050392	050393	050394	050395	050396	050401	050404	050406	050407	050410	050411	050414	050415	050417	050418	050450	050421	050423	050424	050425	050427	050430	050431	5043	5043	043	50043	n
CASE MIX	01.2398	.085	01.2966	.19	.123	01.2459	0104137	.234	01.2615	0104059	01-1616	01.1137	01-1327	01-2004	01010	00.9877	01.2624	01.1035	01.4973	01.1704	01-4667	01.1170	01.1431	91714	00.9989	01-0932	01-1953	01.0569	01.2675	01-2554	01.3236	00.9628	01-2500	01.3389	2041-10	01-0322	01.4418	01.1169	01.2053	00.9911	01.1300	0101247	01+10TO
>	20	50	050299	503	050301	050302	050303	050305	050307	050308	050309	050310	050312	050315	050317	050318	050319	050320	050324	050325	050327	050328	050329	050331	050333	050334	050336	050337	050342	050343	050348	050349	050350	050351	050353	050355	050357	050359	050360	5036	050362	2000	2030
CASE MIX	21	20	21	0	20 1	2 .	7 !	817	208	51	01.2433	V 6	N O	01-1819	(1)	and.	0	01.2936	01.1741	01.4770	01.2452	00.9703	01.1221	01.4550	01.1974	01-3647	01.2448	01.1643	01-1753	67479	00.8658	90	9 1	22	0 0	01.3189	.20	.22	.07	.51		010	OTO
PROVIDER	050228	050229	050530	050231	050232	050233	050234	050235	050236	050238	050239	050540	050242	050243	050245	050248	050251	050253	050254	050256	050258	050260	050261	050262	050263	050267	050268	050269	050270	050273	050274	050276	050277	050278	050209	050281	050282	050283	050286	050289	050500	050202	212000

NOTE: CASE MIX INDEXES DO NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS.

: CASE MIX INDEXES INCLUDE CASES RECEIVED IN HCFA CENTRAL OFFICE THROUGH JUNE. 1987.

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ř	01.2007	100000	01-1611	060050	0101115	070021	01.5389	100018	01-1683
0	01.1815	060004	01.0718	1 40	00.9130	1 00	01.1705	000	01018
050594	32	000000	01.3538	- V	01.1006		01.1692	000	01.15
050597	.128	900090	01.2292	9009	01.0346	0025	01.4699	000	01.37
505	163	00000	01.1298	9009	01.2134	91	01.1936	000	01.1
0505099	71	800090	01.1359	6006	01.2838	0027	01.2532	100024	0101
200	01010000	060000	01.62414	060065	0101326	200	01.3200	100026	010 3
5060	185	010000	01-1690	6006	01-0320	0000	01-1356	100027	010
050605	01.0076	060012	01.3425	060068	01.2651		01.2786	100028	01.1
050607	01.1385	060013	01.2452	060070	01.1590		01.2001	100029	01.1
050608	01.1007	060014	01.3941	060071	01.2487		01.2202	100030	010
506	01.2173	060015	011.3110	0	01.0022		01.2514	100032	01.2
050613	00.9626	910090	01.2166	0	01.0273		01.2684	100033	01.1
050615	01.2029	20090	01.2110	2009	9696 00		01.0602	100034	01.2
020616	01.1244	060018	01.1041	060075	01.2154		01.2088	100035	01.1
050618	1090-10	610090	01.2535	060076	01.2511		01.1353	100036	01.1
619050	01.2376	060020	01.2752	00000	80%600		01.2080	100038	6.10
770056	01.00%	220000	0104300	00000	00.000	400000	0102303	100050	010
000000	4607010	000000	010070	0000	0000140	•	01.0404	10004	
470000	1501010	470000	01.4578	00000	0161310		01.0448	740001	010
020000	01.0716	040024	01.0417	000000	1606-00		11/110	540001	2010
150630	01-1550	050050	01-2279	060060	01-0331	100060	150010	10004	010
050635	01-2069	060028	01.3300	060093	00-8819		01.2553	100046	01-10
050636	01.2203	060029	00.9841	6009	60*6*00		01.3942	100047	01.1
050637	01.1391	060030	01.2049	0	01.0228		01.2009	100048	6.00
050638	00.9210	060031	01.3655	960090	01.1380		01.2285	100049	010
149050	01.0652	060032	01.2984	660090	00.7930		01.1038	100050	01.0
206	01.0119	060033	01.2092	100010	01.5261		0101714	100001	01.0
050644	01.1348	060034	01.2426	000	01.5198	6	01-1301	100052	01.2
506	8797.00	060035	01.2136	010003	01.2106		01.0750	100053	01.1
050649	01.0303	060030	0000000	400000	01.1037		202010	***************************************	100
050650	01-1906	060038	01-2460	000000	01-1762	100001	01-2632	100056	010
150657	01-1063	060039	01-1323	000000	01.1982		00.9658	100057	010
050661	00.9134	140090	01.2405	-	01.1069	10	01.0331	100059	01.2
050663	01-1229	060042	01.0167	-	01.2125	9	01.3258	100060	01.4381
2066	00.5719	060043	01.0571	010010	01.3882		01.5718	190001	01.1
2066	00.8947	440090	01-1618	-	1.186	~	01.2485	00	01.1
050668	01.2266	060045	01.0600	7001	1.176		01.2554	000	01.10
050669	00.8741	940090	01-1028	7001	.197	100010	.187	100065	010
2290672		060047	01.0463	1001	1.126	10001	.988	000	01.1
150673	00.9226	060049	01.1909	070015	76	100012	1.23	0	010
150674	01.1213	060050	01.1831	1001	1.260	100	0.948	100069	010
1906	18	150090	01.2632	1001	1.261	100	·154	0000	01.2
020677	00.8325	060052	01.0294	010018	01.2116	100015	1.10	10001	010
2000	471	060953	00.4522	1001	1.162	100	1100000		010

NOTE: CASE MIX INDEXES DO NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS. : CASE MIX INDEXES INCLUDE CASES RECEIVED IN HCFA CENTRAL OFFICE THROUGH JUNE., 1987.

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CASE M	00.9215	01.0374	00.9323	01.0191	00.9268	01.1792	00.9150	00.9385	01.0534	00.9072	00.9395	6566.00	01.1928	00.9529	01.2003	6790010	000000	00.9040	00.000	01-1589	01-0553	01.2420	00.9162	01.3756	01.0841	01.1116	6266-00	01.6753	01.1887	01-1061	01-1190	00.9086	01.1354	01.1882	01.0056	7410000	01-1515	00-9767	00.9301	00.8959	00.9431	01.0464	00.9332	90%600	01.0543
C ==	(panel)	and :	pod :	pint y	med in	110054	gred	-	pH	pod	and	pri-	-	-	pod g	mel op	nd at	nd pa	of the	of pa	4 (80)	1 000	-	(pad)	-	good is	god s	pool g	200	mel po	d pos		god.	put.	good p	red pe	od pa	d go	d pm	- Bank	- prof	gand	god	pred.	god
136 136	802	122	140	184	000	100-8911	350	989	135	01.1322	666	36	134	01,3131	776	01.0471	170	00.0742	7 1 5	71 2	161	00.9391	101	34	690	184	091	01.0176	111	000	01-1162	986	184	01.0637	239	0 0 0	00.004R	47	001	00.9912	080	385	661	0	5
PROVIDER 100263	2	9	026	02	0 4	020	100273	1100011	110002	110003	110004	110005	110006	110007	110008	110009	110011	110011	110014	110015	110016	110011	110018	110020	110023	110024	110025	110026	110021	110028	110030	110031	110032	110033	110034	110035	110037	110038	110039	110040	110041	110042	110043	110044	110045
CASE MIX 01.1820	1.316	1.179	1.165	10137	1 - 202	1	1.138	1.223	1.124	1.0	1.1	1.2	103	104	100	701	0 1	101	100	0 =	1 0 1	I .O	1.4	1.0	I.I	1.1	0	1.5	I o I	707	8 0	1.1	I . I	1.1	101	107	1 0 1	100	1.2	1.	101	1.01	1.1	101	1.1
PROVIDER 100203	0	0000	100201	01	100209	od mad	1	100213	post	and	021	02	02	100221	20	0 0	100224	NO	220	220	10	100230	CO.	023	023	023	02	100237	02	200	100241	02	02	02	4 .	200	025	20		100255	025	S	025	100260	100262
					20	0101010	382	33	126	01.1666	01.4730	01.1222	00.8808	01.2315	01.0391	01.2880	600000	010-9803	107.10	00-0761	00.9605	01-1489	01.1466	01-1639	01.4374	01-1303	01.1099	01.1472	01.2787	01.0051	01-1846	01.3380	01.2665	01.0764	01-1092	01.0443	01-1175	01-1425	0101749	00.8394	01.0970	01.1863	01.1884	01-2150	01-1406
PROVIDER 100137	100138	100139	100140	00	100143	100145	100146	100147	100149	100150	100151	100152	00	00	00	000	000	000	3 6	50	000	100166	00	00	60	00	00	100173	00	000	000	00	00	00	00		2 5	000	00	00	00	100195	100196	100199	100200
	01.3967		75	2	36	17			08				01.2543	01.1411	01-1212	01.1131	01.2190	00.0487	01 1040	01-0006	635000	01-1707	01.0728	01.1042	01.0022	01-1003	01.1562	00.9341	01.4486	01-1768	9660-10	01.0572	01.0432	9566 00	01.1270	1760-10	01-109	01-2666	01.9451	01.1544	01.1270	01.1482	01.2080	1106.00	01.3906
PROVIDER 100074	100015	07	00	0000	6/0001	100081	0	000	100084	000	100086	0	100088	1000 89	100090	100092	100003	100098	100100	100100	100103	100105	100106	1001001	100108	100109	001	100	100113	100114	100117	00	100120	100121	100122	100	100126		100128	100129	100130	100131	good	10	100135

NOTE: CASE MIX INDEXES DO NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS. : CASE MIX INDEXES INCLUDE CASES RECEIVED IN HCFA CENTRAL OFFICE THROUGH JUNE., 1987.

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375 10	110160	2007 00	130003	01 343	4000	1000	340045	OT JOER
01.2020	110109	00.000	130002	1602010	140007	01-1722	140065	0102033
020	110171	01-1964	130005	0101010	000	01-2154	4006	
596	110172	01-1377	130006	01.5116	4000	0.995	140068	.067
1446	110174	00.8859	130007	.319	10	01,2823	4006	
39	110175	01.0937	130008	00 8935	140011	01-0735	140070	01-1752
0	110176	9011-110	130006	.963	4001	.183	0	.128
48	11011	01.1381	130010	499600	140013	6861010	140074	01.0048
3040	110178	00.9028	130011	01610	10041	103	200	0467910
7676	110101	******	130012	01 1207	1004	201.	140070	000
1001	110183	01-1309	130015	01-1799	4001	141	008	539
47	110184	01-0419	130015	01.0691	4001	01-2657	140081	460°
63	110185	00.9202	130016	00.9346	140019	.950	140082	01.1555
900	110186	01.0390	130017	01.0056	140023	.048	140083	01.0892
9880	110187	00.9760	130018	01.1850	140024	.971	140084	01.1695
150	110188	01-1237	130019	01.0369	140025	01.0836	140085	01.2129
01.0058	110189	9696 00	130021	00.8972	140026	01-1000	140086	01.0360
4110	110190	01.0014	130022	01-1059	140027	01.0675	008	01-1543
.3137	161011	01-1266	130024	01.1630	140029	01.2067	140088	01.3976
00.9034	110192	01.2112	130025	01.0168	140030	01.2848	140089	01.0797
00.8957	110193	01.0622	130026	01-1227	140031	01.0520	140090	01.2087
.9731	110194	01.0177	130027	1668.00	140032	01.0931	140091	01.2999
00.9650	110195	00.9551	130028	01-1293	140033	01.1076	40	01-1130
00.8728	110196	01.3434	130029	01-1282	140034	01.0470		01.0803
1196-00	110198	01.2263	130030	00 63939	140035	4880.10	140095	01.0998
1666-00	102011	01 1705	130031	1024-00	140037	000	140098	01-1765
00.8936	102011	00.9852	130034	00.9421	140038	115	140099	01-1755
01-0208	120001	01-3889	130035	01.0698	140039	.026	140100	01.0650
01-1278	120002	01.0975	130036	01.1190	140040	.093	140101	01-1419
.1545	120003	01.1066	130037	01.0309	140041	978	140102	00.9618
00.8572	120004	01-1596	130038	00.8788	400		140103	01.0966
0	120005	01-1754	130039	01-1444	140043	01.0931	140104	01.0887
0	120006	01.0757	130040	1516.00	140045	01.0554	140105	0102129
9026-00	120007	01.3332	130041	00.8711	140046	01.0857	010	00.9377
+116.	120008	01.0281	130043	348	140041	01.0846	140108	0101315
636	120009	00.9788	400		140048	980	140109	4066-00
378	120010	01.4201	130045		140049	01.2410	140110	01.0785
00.9870	120011	01.2925	130048	1968.00	140051	01.1787	140111	00.9640
0	120012	01.0147	130049	.182	140052	01.1447	140112	01.0751
10	120014	01-1163	130050	01.0712	140053	.390	140113	01.3529
01.0360	120015	31	130051	1.072	140054	1.250	140114	01.1053
25	120016	1.0	130053	.014	140055	.966	140115	01.0430
9038	120018	899	130054	.789	140058	.046	140116	01.2172
93	120019	1.061	130056	37	140059	010.	140117	01.1768
15	120021	1.016	000	1.088	140041	35	140118	.303
•1369	02	1.392	000	017	00	·131	-	1.04
.1886	120024	00.8939	140003	00.9210	140063	01,1516	140120	01.0673
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NOTE: CASE MIX INDEXES 00 NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS.

: CASE MIX INDEXES INCLUDE CASES RECEIVED IN HCFA CENTRAL OFFICE THROUGH JUNE., 1987.

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	212								
	613	4	1.188	023	1.004	00	1014	150071	73
4012	129	M.	1.110	140239	1.321	100	1045	50	65
4012	010	947	1004	+	1.168	150018	1.172	20	53
4012	081	4	1.005	140541	.890	150019	1013	50	30
4012	329	No.	1.198	324	.208	150020	•00	5007	01.1798
4012	176	4	1.250	4	.103	150021	1 . 384	5007	01.0227
4012	043	AL.	1.080		.02	005	1.08	20	01.0993
104		140182	N	A-	55	005		Don I	01.0245
013	122	4	1.067	024	.007	150024	. 124	200	
013	122	NT.	10175	124	.118	150025	.261	200	
013	287	N	1.0	124	.832	005	1176	150082	65
013	183	4	10215	025	· 14	005	.089	200	145
3	133	-a	00.9365	140251	.134	150029	-	5	
01	103	NT.	01.1167	025	01.1987	150030	.134	150085	22
3	166.	M.	9000*10	140253	.130	150031	.975	200	01.0942
013	062	M.	01.1300	025	.220	150032	623	200	36
013	660	- T	01.0862	920	*100	150033	.287	200	01.2134
013	660	-	949164	027	.956	150034	.11	150090	87
014	.01	AL.	01.2206	027	.026	150035		160051	84
5	-8	-	01.0612	140275	.110	150036	.015	150092	01.0253
4	.11	NT.	01,2209	140276	01.8120	150037	141	150094	01.0654
3	.10	A 100	0101010	140280	1410	150038	.165	150095	56
4	.02	-4	01.0337	140281	.349	150039	.036	150096	010010
4	.05	-40	0101989	028	374	150042	.156	150091	01.0674
140146	6 .	No.	01,0302	8	*104	150043	01.1102	150098	01.0486
410	.02	*47	01.1299	8	181	150044	01.1455	150099	01.0779
410	.30	- No.	0101336	140289	01,1958	150045		150100	01.290
4015	e 23	200	01.2600	6	145	150046		M.	01.0373
4015	001	-	01+2375	6	138	150047	.279	יש	01.0197
4015	000	-	01.0412	6	441	150048	.157	R.	01.006
015	0	140211	01.0880	140293	387	150049		150104	C d
4015	910	e .	01.0075	140294	186	150050	• 136	R)	6
4015	600	40.7	01,1113		4 1	150051	0101741	501	01.0250
4010	10		001010	0 0	5 C	250051	1000	0100	77
140159	01:1814	140217	01.1663	- a	212	150054	00-00-0	150110	01-022
4015	4 0		01.0343	0000	1 6	5005	390	5011	0101527
4016	-	-00	01.1208	150002	.216	150057	.011	5011	0101147
4016	0115	140220	01.0541	0	.324	In	1.293	5011	01.0226
0	015	-44	01,3026	150004	186	150059	-	5011	01-1743
4	046	74.	01.2038	150005	137	150060	.10	5012	01.0553
16	.132	*4.	56%6*00	150006	.16	150061	1.110	5012	01.0825
4016	. 924	4022	01.3115	150007	.083	10	1.02	5012	0101396
9104	010	4022	01.0706	000	.264	2	1.129	5012	01.1453
104	00	4023	00.9948	150009	115	150064	-	12	01.6135
4016	.02	405	.175	150010	042	2	1012	5012	01.0880
4017	.07	4023	1.031	120011	.128	900	1.10	5012	01-1356
10	6460	NI	39	150012	29	5	40	150129	60
4017	01.2426		10113	150013	440	00	101	5013	01.202
2107					1	- 日日 日日 日日 日 日 日 日 日 日 日 日 日 日 日 日 日 日		1 11 11	1

NOTE: CASE MIX INDEXES DO NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS. : CASE MIX INDEXES INCLUDE CASES RECEIVED IN HCFA CENTRAL OFFICE THROUGH JUNE., 1987.

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TABLE 3.C HOSPITAL CASE MIX INDEXES FOR DISCHARGES OCCURRING IN FEDERAL FISCAL YEAR 1986

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110011	INDEXES DO NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS.	INDEXES INCLUDE CASES RECEIVED IN HCFA CENTRAL OFFICE THROUGH JUNE 1987.
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PROVIDER 150133	01.1486	160055	01-0675	160111	01.1632	170018	01.0647	170076	01.1164
150134	01-1530	160056	01.0009	160112	01.2366	170019	01.2219	170071	00.962
150135	00.7924	160057	183	160113	01.0559	170020	01-1720	170079	00.892
20136	01.0956	160058	01.4186	110	01.0419	170021	00.9407	170080	00.939
100001	.162	9	202	110	01.0602	170022	01-1336	170081	01.080
20009	01.2252	160060		160116	01-1203	170023	465710	170082	146.00
60000	01-1090	160061	01.0558	160117	01-1116	170025	01.0872	170085	01.015
60009	01.1481	160063	0100797	160119	9446	170026	01.0092	170086	01.406
60007	01.0596	160064	01.2003	160120	00.9082	170027		170087	01.196
80009	01.1351	160065	01.1389	160122	01.0639	170030	110	170088	466.00
60009		9	01.0736	160123		170031	01.0233	170089	00.984
21009	1960-10	160067	01-1450	160124	01-1912	170032	01.0797	170090	01.087
600013	01.2185	160068	01.0992	160126	01.1116	170033	101	170092	01.0687
\$1009	01.0446	160069	01.3094	160129	01.0312	170034	00.9348	170098	00.900
60018	01-0598	160071	01-0132	160131	01-1251	170036	956	170095	010
60020	01.0438	160072	0101273	160132	01.0344	170037	01.1073	170097	010010
60021	660	160073	00.9085	160133	01.1084	170038	1466.00	170098	01.0024
60003	7760-10	160014	00.9862	160134	00.9338	170039	9690°10	170099	9660-10
60024	01.1885	160075	01.0154		00.9790	170040	01.3131	170100	00-8928
	01.4380	160076	01.0011		01.0522	140041	00.9930	170101	01.0297
	01.1168	160077	01.0564	160138	01.0179	170043	2751-10	170102	01.00204
160027	.123	160079	01.2260	160140	61.0275	170044	01.0198	170105	01-2466
60028	1113	160080	2911-10	141091	01.0408	170045	01.0203	170105	966-00
67009	01.2369	160081	01-4569	160142	01-1049	170049	01-1685	170106	01.012
60031	083	160083	01-3620		01.0965	170050	000000	170108	00.938
60032	C1.0245	160085	01.2008	160146	01.2245	170051	00.9779	170109	01.0230
60033	01.2218	160086	9600-10	160147	01.2398	170052	01.0116	170110	00.9175
60034	01.0376	160088	01-0746	161091	01.1606	170053	00.9633	170112	01.012
	01.0658	160089	01-1885	160152	01.0495	170054	00.9943	170113	01.057
60036	01-1378	160090	01.0373	160153	01.3432	170055	1416.00	170114	010027
60037	01.1263	160091	01.1420	170001	01.1313	170051	00.9889	170115	01.153
60038	10000	160091	00-0344	170003	01-0966	170058	33	170117	01.033
	01-1997	160094	01-1067	170004	01.0744	170060	1 10	170119	00.930
60041	01.0597	160095	01.0208	170005	00.9241	170061	01.0817	170120	011.10
60043	01.0423	160091	01.1076	170006	01.1344	170062	337	170121	00.862
	01.2467	160098	01-1323	170007	01.1836	170063	925	170122	01.557
600045	01.3718	160099	-	170008	00.9756	170064	00.9410	170123	01.304
95009	01.0448	101091	.089	170009	01-1089	170066	00.9301	170124	00.933
	01.2734	160102	01.2714	170010	0750010	10001	0180010	170125	000000000000000000000000000000000000000
	611175	160103	2016-00	170011	01.1343	170069	01-0507	170128	00.051
60000	01.0507	160104	01.1445	170012	74	170070	270	170130	01.061
60000	01.0508	160107	01-1150	170015	01-0477	170072	366	170131	01-196
60052			01.1743	170015	01.0491	170073	01.2032	170133	01.225
11.00	010010	,	-0			The state of the s	The state of the s	The second secon	The state of the state of
2007	7631 10	140100	01.0021	170016	01-4128	170071	01-1137	170134	00-00

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PROVIDER	CASE MIX	PROVIDER	CASE MIX		CASE MIX	PROVIDER	CASE MIX	PROVIDER	CASE MIX
170138	01.1561	180028	00.9203		.054	190023		190110	00.9769
170139	1866.00	180029	+90.	60	0.943	9002	01.0764	190111	1.288
170140	00-9941	180030	976	ROI	1-14	9000	150	190112	200
170142	-	180031	020	010	1.098	000	200	190113	200
170143	53	180032	0		179	190029	13	190114	863
170144	01.2780	180033	776	010	1.355	190033	91	190115	210
170145	32	180034	343	010	.239	190034	012	9011	070
170146	01.2504	180035	75	01	0.88	190035	.252	-	.988
170147	01.1278	180036	322	010	875	190036	1.39	96	10.
170148	01.1671	180037	-	0	0.863	190037	9616*00	190119	6.
170150	01-1044	180038	117	110	0.957	190039	01.2821	190120	.940
170151	0966.00	180040	0	011	.952	190040	01.1886	106	-
170152	96	w	00.9787	10	.109	190061	01.3090	9012	.2
170159	86	w	01.0230	011	0.923	190043	01.0784	106	197
170160	00.8967	10	01.0148	011	.987	4006	01.0354	106	0.
170164	01.1605	180044	00.9707	180120	.935	190045	01.0953	190128	7886-00
170166	01.0221	180045	01.1170	01	.03	190046	01.2698	106	9666000
170168	00.9878	180046	01.0060	012	.883	190061	01.0154	106	01.0154
170170	01.1689	180041	01.0037	180123	01.1747	190048	01.0361	190132	01.0264
170171	01.1044	180048		180124	.156	4006	1696.00	190133	01-1339
170172	00.9886	180049		012	806°	190050	00.9832	190134	00.8796
170173	00.8813	w		012	.950	190053	01.0896	106	01.2882
170174	00.9346	w	01.1377	N	.081	9006	01-1764	901	00.9446
170175	01.1036	w		180128	00	190058	00.9755	106	.003
170176	01.2273	w	01.0765	180129	00.9728	9006	01.0668	106	00.7021
170178	00.9032	w	01.0382	PPS.	.184	190060	01-1580	190139	01.0389
180001	01.1326	w		013	0	9006	01.2173	106	1616.00
180002	01.0169	w		180133	01.1251	190065	01.2828	191061	
180004	01.0378	180059	00.9260	MS.	00	9006	01.0270	190142	2616.00
180005	01.0138	~	0	130136		1006	096600	190144	01.0631
180006	00.9234	180062	0.	130137	62	1006	00.8850	190145	01.0130
180007	01.1613	w	1096.00	100061	6.	190075	-	190146	01.3315
180009	01-1002	00	600	190002		190077	01.0349	190147	96
180010	01.4602	w	970	190003	1660-10	190078	01.0124	4	9968-00
180011	9686.00			190004	poted	190079	0	190149	6086-00
180012	01-1337	4	1.04	000	1.130	180061	.935	106	01.0643
180013	01-1255	w	6.0	190006	1.06	190083	00.9461	190152	01-1028
180014	01.4114	w	6 .	000	1.002	8006	260	106	.968
180015	01.0312	w	01.0541	190008	.232	190088	01.0118	190156	0
180016	01.1248	w		000	1.0	190089	01.0425	190157	934
180017	01.1477	w.	6 .	001	0000	060061	01.0571	190158	088
180018	01-1128	w	6.0	001	1.028	190092	01.1034	910	410
180019	01.0916	w.	0.	00	0.93	6006	.928	106	33
180020	00.9526	180081	101	190013	.0	190098	01.3058	5	986
180021	0.0	-	101	100	0.993	600	.045	10	776
180023	.886	180081	1.0	100	1.08	101061	4	0	29
180024	92	180088	1.2	100	1.104	010	.22	910	0
180025	·079	180092	1.0	100	1.06	010	.813	106	. 88
180026	01.0156	180093	91.2004	610061	01.2828	190106	00.9848	190161	01.0223
180027	1.005	180094	96 0	190020	1.04	010	.078	901	.86
NOTE: CASE	MIN TADEVEC DO	I NOT TON O	SOCIAL STO SOL	EDAM DDC	CVCWDT HATTE				

NOTE: CASE MIX INDEXES DO NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS. : CASE MIX INDEXES INCLUDE CASES RECEIVED IN HCFA CENTRAL OFFICE THROUGH JUNE., 1987.

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TABLE 3C HOSPITAL CASE MIX INDEXES FOR DISCHARGES OCCURRING IN FEDERAL FISCAL YEAR 1986

161 123 122	.455 .895 .407 .217	F1224	01.2620 01.1869 01.1258 01.1258	01-1132 01-1697 01-1647 01-0719 01-5771		0 = 8949 0 = 1511 0 = 1511 1 = 1729 1 = 1358 1 = 1358 1 = 1128
10ER C 82 0 83 0	20086 0 20087 0 20088 0 20089 0 20090 0				220119 220120 220121 220121 220122 01 220123 01 220128 01 220128 01 220128 01 220128 01 220128 01 220128	00000000000
CASE MIX 01.1909 01.0847	1,2013 1,1142 1,2450 1,1861	10,0477 10,1735 10,1048 10,1341			006460	3132 3858 54547 3858 17085 1501 1418 139
10ER 22 23	0024 0025 0026 0028	220030 220031 220033 220034 220035	80-289	25222	20068 20068 20068 20068 20068 20066 20066	8 9 0 1 7 5 7 5 9 1 5 9 1
CASE MIX 01.0926 01.1678 01.1733	01.0498 01.2630 01.0336 01.5503	01.1649 01.1108 01.1239 01.2004 01.1379	01.0885 01.1985 01.1732 01.1019 01.1783	01.0221 01.0386 01.1409 01.1879 01.0234 01.1995	01.1260 01.1260 01.3165 01.2307 01.6201 01.1518	01.1044 01.0914 01.1768 01.11768 01.1173 01.2218 01.1397 01.1550 01.2231
œ						220005 220005 220006 220009 220010 220011 220015 220015 220016
CASE MIX 00-9919 01-1851 01-1343	01.0387 01.0777 01.0167 01.1445 01.2281	01-3221 01-1397 01-0560 01-2414	01:0436 00:8824 01:0957 01:1007 01:1206	00.9938 01.0290 00.4590 01.0378 01.2396	01.2508 01.1601 01.2414 01.2414 01.607 01.1605	01.1326 01.18485 01.18930 01.1995 01.21912 01.21889 01.1488
0000	0000	000000000000000000000000000000000000000	00000000			210012 210012 210013 210013 210015 210016 210017 210018 210021 210021 210023
ASE M 1.015 1.118	1.361 1.042 0.976 0.995 1.037	1.0984 1.0985 1.0995 1.083	1.154 1.154 1.154 1.008	0.929 1.061 1.061 1.003 0.933	1.226	HOPEDONFESTOC
8017 9017 9017	9017 9017 9017 9018	010000000000000000000000000000000000000	9018 9019 9019 9019 9019	019900000000000000000000000000000000000	000000000000000000000000000000000000000	0000000000000

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CASE M	.042	.03	.811	017	5.	.059	20	.213	159	AC.	123	16	090	326	152	4	3.3	123	01-1153	01.1036	01.3353	696	186	287	56	4	01.1341	01.1399	72	01.1571	01.0818	34	01.2838	01-1557	01.4954	0	0	661	.328	1.2	1.259	1.58	00.9792	1.09	01.4498	0101622	1.29	1.19	.92	1.22
ROV	400	240009	400	00	4001	1004	240016	4001	240018	240019	4002	4002	240022	240023	01	240025	240026	240027	240028	240029	240030	240031	240033	240036	240037	240038	240040	240041	240043	240044	\$C	40	240047	40	04	35	240051	240052	35	2	3	240057	35	240059	90	4006	4004	4006	900	4006
ASE M	0.807	1.228	10137	1.064	1.20	00	05°	.963	01.1732	86600	0.905	10171	1.136	201	1.064	.043	- part	0194	01.2109		.973	8 46 °	· 258		076		.053	.187	.108	·153	0400	.058	01.0608	1 .028	10144	1.198	1.086	0	10144	013	1.188	1027	10110	1.319	10412	1.167	1.337	0.930	10125	1.073
RO	230203	30	30	3020	0	021	3021	N	3021	3021	3021	302	22	230223	22	230225	2	2	3	3	3	3	230236	33	230238	3023		3024	302	52	3025	3025	230258	305	3026	3026	3026		3027	230273	3027	3027	3027	4000	400	4000	4000	4000	00	400
ASE M	PU!	1460	.085	.29	01.0666	6610	.13	02	1.120	1.133	10114	.39	102	083	1.078	06600	10411	10101	*083	10141	0.872	106.0	.927	359		•167	1.092	10131	1.082	1910	.936	.085	01.0732	.038	0.989	1.023	0012	Iolos	1.089	0.983	06001	0.935	19701	00600	1.146	1.139	1.187	1.088	1.062	64600
ROV	230137	301	301	301	10	301	10	230145	10	410	410	IC	510	10	510	510	015	215	510	315	016	016	910	016		910	317	317	710	110	110	217	017	217	710	018	018	21.0	970	018	010	218	016	610	610	610	610	610	610	020
SEM	.214	140	01.0234	.15	°056	0075	.126	07	.025	440	01.1630	11	01.1809	14	353	01.1068	148	101	388	0	60	61	33	23	01.3299		01.0677	01.1167		00.9510	00.8780	0	20	323	pmd	13	0 0	7 e	5 1	1	000	\$000	97.	270	52	670	151	.079		1.126
IV	3001	3007	3007	3008	3008	3008	3008	3008	3008	3008	3008	3009	3009	3009	3009	3000	3009	3008	3000	3010	3010	3010	3010	3010	30	3010	3010	3010	3011	3011	3011	3011	3011	3011	3011	3011	3011	2100	3012	3012	2016	3012	2106	3012	3012	3013	3013	3013	3013	3013
SE M	01744	.207	0110	0147	.172	.995	e 123	.259	.200	.207	.237	. 186	9460	0057	661°	.168	.277	.520	.057	.985	1610	0043	.500	0110	01.1445	060	660°	.806	0472	.093	9760	.314	0327	090	.958	0400	4084	1110	0110	2000	0000	4110	0010	0160	0214	.106	-	10	-	-1
ROVI	5000	3000	3001	3001	3001	3001	3001	3001	3001	3005	3002	3002	3002	3002	3002	3003	3003	3003	3003	3003	3003	3003	3003	3003	230040	3004	3004	3004	3004	3004	3008	3005	3005	3005	3005	3005	2005	2000	2000	3000	2000	2000	3000	3006	3006	3006	3007	3007	3007	3007

NOTE: CASE MIX INDEXES DO NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS.

: CASE MIX INDEXES INCLUDE CASES RECEIVED IN HCFA CENTRAL OFFICE THROUGH JUNE., 1987.

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3 CHOSPITAL CASE

ROVI	AS	PROVIDER	CASE MIX	PROVIDER		PROVIDER	ASE M	PROVIDER	CASE MIX
S.	1.069	401	0	240187		900	1.046	050	00.930
4007	01.0636	401	0	240192		250043	0-84	250107	00000
4007	.003	401	-	240193		900	1 40-0	250100	057
007	.986	401	01.0681	240194		400	988	250110	00
100	.051	104	4616.00	240195		00	70.07	250111	RTT
007	0147	401	4966.00	240196		700	908	250112	010-0
100	.18	401	01.2593	240200		00	01.2638	250113	1.00
0	01.0016	240132	01.2482	240201	00.9199	2	_	250114	869
100	.31	401	01.0834	240205		500	•	250117	6.0
000	.06	401	01,1346	240206		00	00.8629	250118	043
800	.23	104	90.9144	240207		500		250119	888
800	. 26	401	01,1115	240208		200		250120	0 5
008	.20	401	01.1069	240210		00		250121	656
008	. 25	401	01.0410	250001		250060	00.8364	250122	025
008	.22	104	01.0748	250002		00		N	123
008	.88	401	00.8526	250003		00		N	88
008	.10	401	01.0445	250004		00	- 63	N	085
008	000	401	01,2206	250005		00	- 8	N	392
008	.36	401	00.9993	250006		00		CU	00.8672
008	110	401	01.0338	250007		250067		CV	948
600	.05	104	00.9882	250008		250068	00.8462	M	900
600	960	401	01.0570	250009		250069	- 6	M	01.0489
600	120	401	00.9382	250010		250071		100	930
600	.26	401	00,9393	250012		250072	- 9	100	00.8317
600	10	401	01.0465	250014		250073	164800	250134	01.0195
600	15	104	01.0213	250015		250075		250136	10
600	16.	401	01.0031	250016		250076	- 6	250137	30
600	05	104	00.9945	250017			8606.00	250138	and
010	200	104	01.0200	250018			01.2269	250139	grad .
010	0	104	01.0279	250019			00.8563	250140	00.8622
010	000	+01	01.0485	250020	958		0101119	260001	01.3757
010	910	104	01.0779	250021	00.9756		01-1132	0	200
010	17	4010	01.0195	250023	006		00.8424	0009	01.0158
010	5.	4016	01.1666	250024	956		01.0573	009	01.0125
010	7 1 0	4010	00.9715	250025	646		00.9492	009	01.1266
010	500	+010	01.2483	250026	00.8744	250086		009	01.1766
010	000	4010	2511.10	250027	0946	00	0	09	01.1360
010	200	4010	00.9741	250029	0	250089	00.9451	600	01-1735
110	000	4010	01.0304	250030		250091		00	
110	010	1017	01.0782	250031		250093		1009	
110	000	100	01.0603	200		600	.091	1009	Pro-
110	000	+017	4610.10	2003		600	160	9	22
110	53	de .	01.0150	03	01.1027	250096	0	1009	10
	000	1104	00.9743	5003	6	600	*039	10	366
1104	1.09	101	01.0224	5003		600		1009	334
7104	06.00	104	01.0329	5003	0.913	600	.033	10	180
1104	60	101	01.0027	2003	0.8	010	.180	1009	203
4012	1.006	1018	01.0514	2003	146.0	10	0.86	00	186 0
710	000	101	01.0363	2004	01.0539	250102	1.324	1009	1.007
4016	01.0032	101	1004	20	6.	501		8002	.317

OCCURRING IN FEDERAL FISCAL YEAR 1986 TABLE 3C HOSPITAL CASE MIX INDEXES FOR DISCHARGES

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R CASE MIX 01.0718 01.07589	01.2605	01.1975	00.9673	00.9464	01.2443	01.3945	01.0705	010-2060	01.2299	01.0047	01.0838	01-1520	01.0500	0103702	01.0570	01.0184	01.0831	01.0652	01-1591	01-0085	01-0420	00.9558	01.0397	01-1077	-	01-0523	01-1470	01.2340	01.2177	01.0930	-	10000	01-0416	967	166	366	966	1.072	.103
280017 280018 280020	280021	280023	280025	280026	280029	280030	280031	280032	280034	280035	280037	280038	280039	280041	280042	280043	280045	280046	280047	280048	280050	280051	280082	280054	280055	280057	280058	280060	280061	8008	280063	2000	8008	8008	9	8007	800	280074	8007
CASE MIX 000.9006 000.8757 01.0797	00.9288	BN	273	781	00.8201	1556.00	00.8601	00.9814	01.2785	1926.00	01.1113	00.8576	00.9283	01-1550	01-0454	00.8676	8988	00.9307	336	6 4	34	-107		63	00.6897	050	1856.00	00.9150	01.1181	01.0793	9775010	0200010	01.2169	01.2197	1.07	1023	1.368	0.96	96
PROVIDER 270030 270031 270032	270033	270036	270040	270042	31		270046	- 00	-	0		250052		1		6	0		270061	0 -	2	3	4	270075			. 18					ROOOS	80008	80010	80011	80012	80013	80014	0015
- Per 1000 1000	.073	01.0545	m -	01-1621	10	PO 1	A 10		-	-		~	00.9739	-	01-1825	00.7855	01-1357	01.0932	1180010	01.0487	01-1145	01.5478	.067	8106-00	889	01.0552	.269	.157	154451		0.989	.899	.007		1000	.888	1776.	1.	00.9463
80V 601 601 601	910	00	0171		5																				270009		270012		270014						70024		70027	0028	10029
SE. 2	1.057	01.1039	1.244	013	1.0	01.0514		01-1328	01-1532	00.9913	01.1964	01.6617	01-1751	01.4754	1956.00	01.2486	01.0508	01.2090	010010	01-1123	01-1752	01.1533	01.1230	01-0724	01.0209	01.0765	00.9881	01.1723	01.0260		1768	5237	5798	1376	2650	1918	1.0	96	01.1020
PROVIDER 260081 260082 260083	08	08	60	260092	260093	260094	260099	260097	260100	260102	260103	260105	260107	260108	560109	260110	260111	260112	260115	260116	260118	260119	250120	260122	260123	260127	260128	260129	260131	260134	260137	260138	260141	241092	4109	4	109	14	6015
E M 232 191 191 160		.283	01.0932	.402		01.0205	00.9670		01-1487	01.0941	0 0	. 0	01.0954	8766.00	01-1663	01.1588	00.8436	01-0485	01-1262	01.0828	01.1830	01.0384	01-1269	01.2057	00.9841	01.0872	01-1284	01.1636	01.1026	326		01.6489	.003	996°	0	01.1917	.061	•00	00.9953
260021 260022 260022 260023	NI OL C	260027	260029	260031	260032	400	260035	009	260037	660002	260041	260042	260044	260045	260047	240048	260050	260051	1 10	260053	6009	550055	260057	260058		260061	260062	260065	260065	260066	260067	260068		260073	260074	260077	P- 1	260079	m

NOTE: CASE MIX INDEXES DO NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS. : CASE MIX INDEXES INCLUDE CASES RECEIVED IN HCFA CENTRAL DFFICE THROUGH JUNE., 1987.

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OCCURRING IN FEDERAL FISCAL YEAR 1986

TABLE 3C HOSPITAL CASE MIX INDEXES FOR DISCHARGES

NOTE:

	290013	01-0417	310000	01-1107	210066	CASE HIA	PRUVIDER	CASE HIN
01.2169	200013	01-0502	310010	01 1750	310064	07/10/10	320016	01.00996
01-0142	290015	00.0160	310011		310060	0101/8/	320018	01-1417
0320	290015	01.0576	210012	01 1000	310060	010000000000000000000000000000000000000	220025	1147010
9683	290018	00-9070	310012	136	310070	010001	320021	01-1076
2485	290919	01.1576	310014	20.	310071	4 4	320025	01.0325
9151	290020	00.9120	310015	01.2167	310072	01-1408	320030	01-0080
01.0721	290021	01.4080	310016		310073	01-1685	320031	00-9142
.9568	290052	01.4370	310017	01-1509	310074	-	320032	00.9877
01-1856	290027	00.9530	310018	01.0995	310075	01-1915	320033	01,1838
01.2498	290028	00.9719	310019	01.4337	310076	01.1958	320035	00.9223
01.3672	290029	00-8045	310020	01.1220	310077	01.3710	320037	14
00.9263	290031	01.2186	310021	01.1594	310078	01-1248	320038	01.0933
00.9759	280062	01.2412	310022	01-1553	310081	01.1593	320046	01.0704
01.0422	290033	01.0776	310024	01-1422	310083	01-1276	320048	01.0562
•9394	290034	01.0036	good	01.1053	310084	01.1783	320049	01.0546
8068 00	300001	01-1828	310026	01.1278	310085	01.1674	320051	01.0554
6110-10	300008	01.0869	and .	01.1643	310086	01.1762	320053	1196.00
00.9982	300003	01.4949	310028	01.0953	310087	01-1421	320056	00.8646
866600	30000€	01.1991	310029		310088	01-1220	320057	01-1178
01.0400	300008	01.0630	310031	02.2514	310090	01.1607	320058	00.8422
6266 00	300007	01.0782	310032	01.0896	310091	01.1656	320059	01-1098
0166.00	300008	01.0748	310033	01.0601	310092	0101648	320060	00-9590
00.9825	300009	01.0932	310034	01.1231	310093	01.0444	320061	01.0767
01.1387	300010	01.1282	310036	01.1529	310094	01.0874	320062	00-8924
01.0231	300011	01.1424	310037	01.1699	310096	01.2858	320063	01-1272
9446 00	300012	01.1575	310038		310105	01.0427	320065	01.0982
	300013	01.0604	310039	0101935	310108	01.1760	320066	00.8221
00.9720	300014	01,1839	310040	01.1249	310110	01.1592	320067	9006-00
01.0574	300015	01.0810	310041	01.1852	310111	01-1709	320068	01-1283
01.0974	300016	01.0152	400	01.0957	310112	01-1396	320069	6160010
01.0228	300017	01.2445	310043	.133	310113	01-1284	320070	01.0834
	300018	01.1685	310044	01.1634	310115	0911-10	320071	1146-00
01.0038	300019	01.0844	310045	01-1470	310116	01-1992	320073	00.9628
	300020	01.1180	310047	01.2640	310118	01-1330	320074	01-1823
	300021	01.0654	310048	·154	310119	01-1523	320075	00.5716
00 8 8 6 6 4	300052	01.0874	310049	.137	310120	01.0623	320076	01.0848
00.8682	300023	01-1240	310050	.104	320001	• 307	320077	9066-00
00.6412	300024	01.1917	310051	.236	320002	01.1820	330001	01-1371
	300028	01.0929	310052	.174	320003	01.1598	330002	1611-10
00.9284	300029	1.	00	.139	320004	01.0948	330003	01.2228
. 3360	300032	920	310054	1 . 254	320005	01.1859	330004	01-1092
01.1660	300033	770	100	.194	320006	01.1463	330005	01.3672
.9953	300034	01.2095	310057	01.2125	320009	01.2114	330006	01.2261
•3906	310001	1.36	500	1.012	320010	01.1867	330007	01.1326
•1794	310002	01.6083	310059	00.8322	320011	01.0235	330008	01-1123
.2872	310003	01.1217	310060	01.2169	320012	9666 00	330009	01.0084
.4100.	310005	1.0	310061	0101010	320013	966.0	3001	01-1127
06860	310006	7771	310062	00.0751	320016	-	2000	0761 10
		9			*1007C	666600		

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E C	4.026	Les S	1 0 0 0	1.069	1.160	1.22	loc	Dol		-	00	62	00	01.1085	90	6.0		- C	T 200	33	060	885	342	00.8246	01.4927	01.0071	162	983	884	e779	.766	70	773	.763	.859	122	856	610	0113	9 70	890	1.602	Loll7	1.548	
IVO	200	3029	3 6	3029	3030	3030	3030	3030	3030	3031	3031	303	303	303	303	303	200	0000	のでは	303	303	303	303	303	303	330351	303	303	303	303	3036	3030	3036	3036	3037	30	3037	3038	00000	3038	3038	3038	303	3039	330393
CASE MIX	200	200	10	19	766	15	60	132	153	361	190	12	15	90	57	70	200	200	5	5	380	151	547	29	36	503	003	120	128	60	77	0 0	157	96	38	74	0	73	ח מ	020	14	30	60	40	40
ď																										330250																			
SE	174	147	245	016	.045	.209	.071	.286	-	030	315	.056	.200	189	040	2011	046	073	01-1558	172	156	278	155	01.1235	440	00.6658	130	,224	362	346	100	188	129	220	134	272	171	126	107	040	082	10	015	507	132
ROVIDER	30167	30158	30159	30160	30161	30162	30163	30164	30165	30166	30167	30168	30169	30171	41100	20100	30177	10179	30180	30181	30182	30183	10184	30185	00100	330189	16101	10193	10194	10195	06100	86108	66101	10201	10202	10203	+0200	20202	0200	0210	10211	10212	10213	10214	10215
SE M	1 - 1 64	00.99	1.289	127	1,168	10422	10147	1.0041	10176	10118	1.024	1.0	00.6214	01.3671	77070	01-1610	01.3999	01.1644	01.1930	6506.00	00.9692	01.1136	00.9788	01.0797	01.0000	01.2436	01.2096	01.4357	01.0296	01.0963	01.1038	01.1088	01.1584	01.0974	01.1032	01. 1361	1077810	01.1279	01.1636	01.0319	01.0314	00.9022	01.0658	01.2185	01.1622
PROVIDER													30100	10105	30102	30104	30106	30107	30108	30109	30110	30111	30114	30115	30110	30118	90119	30120	30121	30122	30126	30127	30128	30132	30133	20135	07100	30141	30142	30144	30148	30150	30151	30152	30153
CASE MIX	53	1013	.006	.130	660	003	52	8448	-	0175	6173	and C	0710.10	01.1990	01-0363	01.0772	01.0536	01.0780	9450-10	01.1073	01.1673	01.1502	0101996	01.1238	01-2098	01.0374	01.0838	01.0429	01.2121	01-1753	31	32	01.1742	.960	.186	2011-10	216	233	.09	0149	1.04	-	1.08		1051
PROVIDER 330013	*	300	01	30019	30020	022	30023	30024	0025	30027	30028	6200	30033	•			38	30039	30041	~	30044	10	9400		30040	30050	30052	53	055	30050	30058	30059	30061	30062	30064		30067	30072	30073	30074	30075	30076	30077	8100	

NOTE: CASE MIX INDEXES DO NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS. : CASE MIX INDEXES INCLUDE CASES RECEIVED IN HCFA CENTRAL OFFICE THROUGH JUNE. 1987.

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OCCURRING IN FEDERAL FISCAL YEAR 1986

TABLE 3C HOSPITAL CASE MIX INDEXES FOR DISCHARGES

330395 01-1701 330396 01-0547 330397 01-0528 330399 01-0322 340001 01-1607 340002 01-6377 340005 01-2821 340005 01-1657 340007 01-1657 340007 01-1657 340007 01-1657 340010 01-2817 340010 01-2818 340011 01-2818 340012 01-1717 340014 01-2516 340015 01-1614 340019 01-1614 340020 01-1024 340020 01-1024 340020 01-1270 340020 01-1270	340050 340051 340052 340055 340055 340060 340061 340060 340070 340071 340071 340071 340072 340072 340073 340076 340086	01-1104 01-1639 00-9794 01-3269 01-3269 01-1540 01-1140 01-1133 01-0265 01-1563 01-2652 01-0923 01-0923 01-0755	340121 340122 340123 340124 340129 340129 340131 340132 340132 340133 340133 340133	01.0542 01.1299 01.1299 01.1733 01.1733 01.1733 01.1236 01.1626 01.2304 01.1861 01.0125 00.9331	350015 350016 350017 350020 350021 350024 350027 350027 350027	01-4091 01-0977 01-1322 01-0106 01-3025 01-1930 01-0183	6001 6001 6001 6001	01.2099 01.0992 01.1134 01.3379
	555 555 555 555 555 555 555 555 555 55	01.1639 00.9794 01.3269 01.3269 01.1540 01.1140 01.1133 01.0706 01.0706 01.0706 01.0706 01.0706 01.0706 01.0706 01.0706 01.0708	340122 340123 340124 340126 340129 340131 340132 340135 340138 340138 340138	01.1299 01.1265 00.9745 01.1733 01.1733 01.1733 01.1236 01.1626 01.2304 01.1861 01.0125 00.9331	350016 350017 350019 350020 350024 350024 350027 350027 350027	.097 .010 .302 .193 .018	360013	01.0992 01.1134 01.3379
	555 555 555 555 555 555 555 555 555 55	000.9794 01.3269 010.9743 010.1540 010.1140 010.1133 010.1047 010.0216 010.0216 010.0655 010.0655 010.0655	340123 340124 340125 340127 340127 340133 340133 340135 340135 340137 340137	0101265 0009745 0101733 0101733 0101236 0101236 0101236 0101236 010125 0009331	350017 350018 350020 350021 350024 350024 350027 350027 350027	132 010 302 193 018	360014	01.1134 01.3379 01.2076
	884 65 8 8 9 9 8 9 9 8 9 9 9 9 9 9 9 9 9 9 9	01.3269 00.9743 00.9743 01.1140 01.1140 01.0133 01.0133 01.0216 01.0216 01.0216 01.0253 01.0455 01.0455	340124 340125 340127 340127 340129 340130 340131 340135 340137 340137 340141	00.9745 01.3120 01.1733 01.1236 01.1070 01.2304 01.21861 00.9331	350018 350019 350021 350024 350024 350027 350027 350027	.010 .302 .193 .018	1009	01-3379
	5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5	00.9743 01.1540 01.1140 01.1133 01.1133 01.0706 01.0216 01.0563 01.563 01.5655 01.0655 01.0765	340125 340126 340127 340127 340131 340133 340135 340135 340136 340137	01.3120 01.1733 01.1236 01.1070 01.1626 01.2304 01.0125 00.95331	350019 350020 350021 350024 350027 350027 350027	.302 .193 .018	-	01.2076
	665 665 665 665 665 67 67 67 67 67 67 67 67 67 67 67 67 67	01.1540 01.1140 01.1133 01.1133 01.00706 01.00706 01.00716 01.00716 01.00713 01.1738	340126 340127 340129 340130 340133 340135 340136 340136 340141	01.1733 01.1236 01.1070 01.1626 01.2304 01.1861 00.9331	350020 350021 350024 350024 350027 350027 350027	0193	1009	
	000 665 777 777 880 884 884	01-1140 01-4685 01-1133 01-0706 01-0216 01-1563 01-5143 01-2655 01-0923 01-1738	340127 340129 340130 340133 340133 340135 340137 340137	01-1236 01-1070 01-1626 01-1861 01-0125 00-9331	350024 350024 350025 350027 350027 350029	0100	09	.270
	666 666 666 666 666 666 666 666 666 66	01-1133 01-1133 01-0706 01-0216 01-1563 01-2652 01-0923 01-1738	340130 340131 340132 340133 340135 340136 340141 340141	01.1626 01.1861 01.0125 00.9331 00.9573	350024 350025 350027 350029 350030	100	1009	01.1809
	665 665 665 665 77 77 77 77 77 77 77 77 77 77 77 77 77	01-0706 01-0706 01-0216 01-1563 01-563 01-2652 01-0923 01-1738	340131 340133 340135 340135 340137 340141 340141	01-2304 01-1861 01-0125 00-9331 00-9573	350025 350027 350027 350029	C	360019	1441010
	655 668 77 77 77 77 78 88 88 88 88	01.1047 01.0216 01.1563 01.563 01.2652 01.0455 01.1738 01.1274	340132 340133 340135 340136 340131 340141	01-1861 01-0125 00-9331 00-9573	350027 350029 350030	277	360021	2410
	668 669 772 775 775 886 886	01.0216 01.1563 01.5143 01.2652 01.0455 01.0738 01.1274	340133 340135 340136 340137 340141 340142	01.0125 00.9331 00.9573	002	.957	360022	.063
	668 669 777 777 775 779 884	01.1563 01.5143 01.2652 01.0455 01.0923 01.1274	340135 340136 340137 340141 340142	00.9331	350030	00.9260	360024	01.0931
	669 772 773 775 775 886 886 887	01.5143 01.2652 01.00455 01.1738 01.1274	340136 340137 340138 340141 340142	00.9573		072	360025	
	777 777 775 775 775 880 884	01.2652 01.0455 01.0923 01.1738 01.1274	340137 340141 340141 340142	01-0416	350031	00.9662	6002	01-1322
	772 773 779 880 884	01.0923 01.0923 01.1738 01.1274	340141	0410	350032	01-1277	360027	01.3446
	27.5 27.5 37.5 38.4 88.5 7.5 7.5 7.5 7.5 7.5 7.5 7.5 7.5 7.5 7	01-1738 01-1274 01-0523	+ +	61 505 10	350033		360028	01010
	775 775 886 885	01-1274	۲	01-1022	350035	00 0730	360020	01.0492
	75 880 885 78	01-0523	3	01-2760	350036	00.00	360031	01-1779
	884 885 87	777	. 4	01-1324	350038	326	360032	01-1270
	84	00.9619	*	01.1015	350039	.973	360034	01.0780
	884	01.0062	*	01.0035	350041	4916-00	360035	01.3109
	85	01.0851	\$	01.1821	900	995600	360036	01.1007
	1 1	01.2296	\$	01-1933	350043	01-1548	360037	01-4590
	-00	01.0620	340151	9690-10	350044	1406.00	360038	01.1986
	200	01.1125	340153	01.8688	350047	00.9521	360039	01.1437
		01-1269	340154	00.3179	350048	00.9503	360040	01.1503
	91	01-4546	340156	00.9015	350050	00.0305	340062	01-0512
	93	01.0651	340157	01-1928	350051	.883	360044	01.1408
01.1	46	01-1913	340158	01.0746	350053	9884	360045	01.2892
01.5	96	01-1122	340159	01.0753	350055	.877	6009	01.0163
01.5	16	01.0221	340160	01.0846	350056	00.9237	360047	01.0120
	98	01.4511	340162	01.1813	350058	00.8834	360048	.354
26.4	66	01.2044	340164	01-1543	0	01.0306	4009	01-1185
010		011.1924	340166	01.1808	350061	01.010	360050	01.0929
01-088	100	00.0273	350003	01 3771	350064	0000	360051	6007.10
6 01.019	05	01-2291	350003	11.0831	350065	010	o v	01-1806
7 01.196	90	01.0508	350004	01.4686	5006	00.9868	6005	0101579
8		0	350005	01.1659	5006	.847	360055	01.1878
1	60	N	350006	01.0750			6009	.139
0 01	11	C	0	00.9290		01-1205	9009	01.0313
_ ,	12	• 074	2000	2668-00	360003	.224	360058	.064
7 7	13	• 705	350009	•15	0	-401	9009	.24
340044 01.0241	340114	No	350010	0.5		0.	9009	1.019
100	71	0670	110000		300000	torer.	0000	0.883
6.10		7804-10	350012	6100010	360009	01.1716	360062	01.3355

R CASE MIX	01-1202	00.9922		38	01.3985	D 4	01-1923	1 00	6	00.9728	1766.00	01.8034	01.2327	00.9717	00-9405	00.9593	00.9617	01.0236	01-4294	01.0690	01.1783	01.1453	00.9340	01.0600	00.9513	00.9862	01.0122	99955	01.0110	0100136	01.2057	01.0339	01.2240	.177	01.0529	100010	00.00	00.9983	01.1698	01-1272	00.9193	000000
ш	370089	370090	0	370092	370093	370005	370096	370097	370099	370100	370103	370105	370106	370107	370108	370110	370112	370113	370114	370117	370121	370123	370125	370126	370130	370133	370136	370138	370139	370140	370144	370146	370148	370149	370153	370154	370157	370158	370159	370161	370163	370165
CASE MIX	01.1582	0660	.152	•135	01.1993	01.6102	040	.066	01.1649	01.0207	01.0754	1.347	754	01.5105	01-1465	01.0810	1600-10	00.8865	90%6-00	01.0434	010-0104	00.9714	01.1227	01.0236	000.9600	01-1699	01.1775	01.0751	.087	00.9739	00.9340	01.0797	00.9740	.900	00.9172	01-1328	01-2948	6160	01.0309	1086-00		00.9224
PROVIDER	370020	002	370022	370023	370025	370028	370029	370030	370032	370033	370034	370035	370036	370037	370039	370040	370041	370042	370043	370045	370047	370048	370049	370050	370051				370060	370063	370064	370065	370069	370071	370072	370075	370078		370080	370082	370083	370084
SE M	01.2025	.048	01.1027	01.1504	01.8290	01-1258	00.8847	01.1326	01.0326	01.0188	01.2004	01.1448	01.0587	01-1235	01-0991	01.1003	01.1031	01.0797	01.0717	01.2111	01-1454	01.1983	01.0679	01-1725	0101010	01.0180	01-1263	01.1070	00.9076	01-2223	01-0071	01.0563	00.9311	01.1222	01.1434	00-9670	00.9516	01.3452	01.0444	01-1032	·19	6866 00
PROVIDER	360176	360177	18	61	360180	40	98	87	60188	68109	35	93					350204					0	360231	01	360234	0 00		0		350345					370007		370012	370013	370014	370015	370016	370017
CASE MIX	01.0708	1.1	01-1474	8071-10	01-1036	01-125	01.0102	117	00.9837	01.0917	01.1144	01.1148	01.2397	01.3136	00.9796	01.3812	01.0722	01.0633	01.2463	01.0324	01.2066	01.2563	01.0966	01-1220	01-1924	01.2553	01.2405	01.0976	01.0471	01-1031	01.0929	01-1052	01.0894	01.4014	01.0526	01.0015	00.9014	967	958	950	0	01-1590
PROVIDER	360119	360121	360122	571095	360124	360126		360128	360129	360130	360131	360132	360133	360134	360136	360137		360140	360141	340145	360144	360145	360147	848		21	52	53	360154	360156	360159	360161	360162	360163	360164	360166	360167	360168		360170		360172
CASE MIX	01.2911	01-1711	01-1374	01-1038	01-0298	01-1843	01.1800	01-1506	01.1836	01.2203	01-1558	01.2427	01.0978	01.3241	01.1106	01.1818	01.0662	01.2543	01-4100	01.1311	01.0541	01.1131	01-1246	01.1818	01.0517	01.0945	01-1595	01.0774	01.2290	01-1754	01.3673	01.2107	01-1882	01.0851	01-1737	01-0690	01-0193		01-3203	177	•003	01-1232
EK							360071				9	1	m (360081	~	3	+			360088			_ ,		360094		9	200	360100			360103			. «		0	~	e .	* 1	360115

NOTE: CASE MIX INDEXES DO NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS. : CASE MIX INDEXES INCLUDE CASES RECEIVED IN HCFA CENTRAL OFFICE THROUGH JUNE., 1987.

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12	7690-10	ROZ	01.0275	200000	1003	SOONS	CASE MEX	PROVIDER	
1000	0 0	000	00000	010066	7901010	340063	1.0390		01.0281
	9 (800	6	390011	8	390064	10141	390119	01.2154
101	· U 38	800	0	390012	N	390068	01.1726	390121	01-132
101	939	800	2	1006	10	9006	01.1755	390122	01.067
101	0163	800	338	1006	016	390061	01.3492	390123	01.1520
101	9 B 4 4	800	01-1026	390015	073	390068	.129	390125	01.105
101	1710	800	253	390016	090	390066	01.1779	390126	01.164
101	.986	800	6	390017	0	390070	200	390127	01.065
7017	.041	800	good	390018	145	390071	01.0743	390128	1110
7017	.979	800	.248	390019	.072	390072	00.9867	390130	01.052
7018	0016	800	01.1451	390020	01.0770	390073	1.12	390131	01-156
7018	.936	8	01.2027	390021	1611010	390074		m	01.0554
7018	.965	800	00.9719	390022	01.1107	390075	.17	60	01-2467
7018	.036	800	01.0154	390023	01.1145	390076	.157	(1)	01-1252
7018	.274	800	01.2724	390024	00.7516	390077	01.1219	100	60
7018	. 709	380061	01,3881	390025	00.7566	390078	.036	390137	01.0876
8000	.332	8008	00.9072	390026	C 3	390079	1.552	390138	01-238
8000	.162	8008	01.1466	390027	01.2946	390080		9013	01.35
8000	.194	9008	01.1538	390028	01.4168	390081	01.2084	9014	01.355
8000	602°	380065	01.1242	390029	01.2828	390083	01.1049	390143	00-00
8000	169	8006	01.2037	390030	01.0772	390084	00.9787	390145	01-10
8000	1117	8008	01.0627	390031	01,1251	390086	01.0587	390146	01-003
8000	439	80	00.9760	390032	01,1322	390087	01.0168	390147	01-08
8000	.076	380070	01.0115	390034	01.1087	390088	216	390168	01.08
8000	0440	80	01.1545	390035	01.1613	390090	01.3439	390149	0101337
8001	0173		6056 00	390036	01,1615	390091	01-0647	390150	01.07
8001	1111		00*6*00	390037	01.1493	390092	01.0435	390151	01-1646
3001	104		1661*10	390039	01.0198	390093	01.0858	390152	01.08
8001	.146		00.9434	390040	989	390095	660°	390153	01-163
3001	918		01.0257	390041	01.1003	390096		390154	01.08
8001	510		01.1963	390042	01.1174	390097	01.2135	100	01.2377
3001	420		0096 00	390043	01.0017	390098	01.3995	10	010173
3001	101		01-1737	390044	01.3829	390099	01.0832	390157	01.1156
3005	201		01.0753	390045		390100	01.4350	390158	01.2053
3005	225		01.2873	390046	01.2939	390101	01.1690	10	01-168
3002	126		526	390047	01.3471	390102	01.2137	390160	01.052
3002	092		020	390048	01.1261	390103	01.0408	390161	00.9925
3002	153		01.2522	390049	01.2519	390104		390162	
3002	283		01.2701	330050	01.1332	390106	.033	390163	01.1361
3008	155		01.2146	390051	01.5917	390107	.119	390164	01.4633
3002	120		00.9308	390052	01.0772	390108		390165	01.0511
3002	039		01.1830	390054	011150	390109		390166	01.0855
3003	081		60	390055	01.4676	390110		390167	01-12
3003	960		01.1245	390056	.104	390111	1.524	390168	01.0835
3003	393		00	390057	01.2597	390112	9600	9106	01,1077
3003	,209		01.0113	390058	01,1961	390113	-103	100	01-5537
3003	064		.386	390059	10	390114	-073	100	
300	01.0881	390007	1.10	9006	137	390115		390172	01-045
3003	200		1-09	3 (203	300116	1.110	- 1	CCCOOL TO
3003	204		0000	100046	020	270110	41101	570113	THE PARTY OF

NOTE: CASE MIX INDEXES DO NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS.

: CASE MIX INDEXES INCLUDE CASES RECEIVED IN HCFA CENTRAL OFFICE THROUGH JUNE. 1987.

PROVIDER	CASE MIX	PROVIDER	CASE MIX	PROVIDER	CA	PROVIDER	CASE MIX
4206	800	200	50	420084	e 70	430060	00.9830
9024	01.0624	420026	210	420085	164	430062	00.8132
9024	-	200	132	420087	.289	430065	992
9024	1.01	NI	8 46	420088	01.1333	430066	00.8939
3900354	01.0544	420029	105	420089	·107	430072	01-0252
06	01.1297	n m	" MI	430001	01.0585	430073	01.0243
CU	11	420032	352	430005	01.0833	430077	01-1959
N	29	(25)	118	430007	01.0122	430079	6
902	01.1549	420035	999	430008	01.1292	430080	00.8924
902	01.3163	2003	447	430009	01.0120	430081	00.9287
902	01.1601	20037	01.1018	430010	01.0503	430082	06.
1 6	01.1130	20038	01.0535	430011	01.1722	430083	. 8
706	01.0636	20039	01.0398	430012	01.2496	430084	856
90208	01.1130	20040	01.1857	430013	01.1081	430085	00.8919
10	01.01413	20042	01.0459	430014	01-1711	430086	01.0052
717	1788 000	20043	01.0459	430015	01.0567	430087	00.8621
010	00.8183	2007	01.0703	430016	.277	430088	8466-00
100	0101/3/		01.0357	430017	9	100044	01.0801
200	01.1035		5166.00	430018	956	440005	01.3283
25	01-2479		000000	430020	0000	440003	01-1074
90001	01-1965	420053	000,9579	430023	01-0114	440005	00.9616
100	01.3388		01.0518	430024	00-9802	440007	00-000
80001	01.1001	20055	00.9811	3002	00.9882	440008	100
600	01.1721	.0	01.0612	OI	96 96 74	600044	01.0134
010	00.9479	20057	01.0615	430027	0104614	440010	01.0035
11	01.1368	20059	01.0945	430028	00.9451	440011	01.1472
71	01.3541	20061	01.0574	430029	00.8938	440012	01-1500
	01.0987	29002	01.0549	430030	01.0819	440014	00.9827
410014	01.1330	20064	01.0245	430031	00.8746	440015	01.4384
	01.0059	20065	01.1511	430033	01.0972	440016	00.9530
000	0101243	20000	00.9295	430034	01.0024	440017	01.2475
420004	000000000000000000000000000000000000000	190074	4710010	430036	01.0265	440018	01.1137
200	010 3500	20070	01 0234	430037	00.8825	610044	01-3086
2000	01-1302	60002	0100000	430038		440020	2066-00
200	01.2711	2002	0101010	5000	1716.00	440022	01.0533
200	717710	11007	010000	+ .	00.9885	440053	99600
200	01-0100	27000	0160000	140064 730064	7600010	440054	01-1232
200	01.0338	2002	0010000	430042	01.0420	440025	01.0769
200	00.0035	1000	0 0	430043	01.1026	440026	01.1095
200) =	2004	0740	420044	0 0	440028	01.0182
200	01-0298	2007	270	43004	01.0028	670044	01.1538
420017	0.9	2007	273		0 0	440030	01.0986
200	I.	2008	050	430057	0 0	150044	186.00
420019	1 000	2000	1000	10000	0000	750044	10010
200	0000	190074	100	G	00 00 C	220077	400
420020	1.063	420081		430054	00.9085	440033	01.0352

NOTE: CASE MIX INDEXES DO NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS.

: CASE MIX INDEXES INCLUDE CASES RECEIVED IN HCFA CENTRAL OFFICE THROUGH JUNE., 1987.

TABLE 3C H	TABLE 3C HOSPITAL CASE MIX INDEXES	MIX INDEXES	FOR DISCHARGES	ES OCCURRING	Z	FEDERAL FISCAL YE	YEAR 1986	Page 2	0 of 24
PROVIDER	CASE MIX	PROVIDER	S	3	CASE MIX	PROVIDER	CASE MIX	ROVI	CASE M
440038	1266.00	440113	1.02	83	01.2042	15	-	0111	
440039	01.3993	440114	01.0380	440184	01,1003	450046	01,1995	011	
040044	916	440115	.065	01	01.0716	450047	01.0326	011	
440041	00.9225	440117	.861	0 2	01.0012	3	01.0691	110	
	01.0365	440120	- (440187	01.0145	450050	01.0254	and s	01.2182
4004	3 - 4	440121	01.0263	440189	01.2136	2000	01.5088	TIO C	0102210
840044	01.3140	440175	28870	440191	2666.00	D U	01.012.0	450113	0101933
440044		440120	01-1000	440192	01-0255		01 - 1805	400	010-0187
440051	98	440131	01-0231	440194	01-0632	5005	0100100	450123	01-1151
440052	918	440132	00.9928	440196	00.9980	450056	01.3220	450124	01.3812
440053	15	440133	01.3023	440197	01-1853	5005	01.1535	450126	01.2435
440054	00.9539	440134	01.0861	440200	00.9888	450058	01.2749	450127	01.0339
440055	01-1123	440135	01.0832	440202	00.9003	450059	01-1460	450128	01-1547
440056	00.9498	440136	01.0705	440203	01.0564	5006	01-1988	501	01.2904
440057	00.9775	440137	01.0278	440204	01.2081	2006	9776 00	501	01.1554
440058	01.0184	440141	00.9161	450002	01.1433	2006	01.3226	501	01.3092
440029	01.0702	440145	00.8380	420004	01.0793	450065	01.0346	201	CI.1714
440060	01.0571	440143	00.9543	450005	00.9567	5006	01.3408	105	01.0865
190044	0101166	440144	01.0542	450001	01.2053	5006	01.2702	201	0103452
440063	01.0881	440145	00.9650	450008	01.1768	5006	00.9853	501	00.7978
440064	00.9207	440149	00.8925	450010	01.1610	5007	01.0954	501	01.1869
440065	01.0409	440147	00.7415	450011	01.2187	5007	01.1340	501	00.8516
440067	01.0276	440148	01.0764	450013	01.2103	450073	01.0041	201	00.9160
4000	01.1276	641044	01.0089	420014	01.0091	5007	01.0788	501	0101245
	01.1337	440150	01.1753	450015	01.3143	450077	01.0221	501	01.0341
0/0044	00.9587	440151	01.0613	450016	01.2840	1000	8866.00	200	01.0369
440072	01-1223	440122	00.0054	450010	101010	420080	01-1413	450146	200000
440073	01-1442	440184	00.8018	450020	017-020	2000	01-1815	300	01-1562
440074	00.9141	440154	01-2091	450020	01-6930	450082	00.9422	450148	01-1550
440078	00.9401	440157	00.8963	450022	01.0187	5008	01.2652	501	01.1834
440079	00.8716	440159	01.0906	450023	01.2998	450084	01.0786	450150	01.0531
440081	4960*10	440160	00.9616	450054	01.1736	5008	01.0934	501	01.0239
440082	019910	440161	01.4237	450025	01,2815	5008	01.2186	450152	01.1973
440083	00.9403	440162	8066-00	450027	01.1044	5009	0101349	501	01.2731
440084	01.0410	440166	01.1936	450028	01.2483	2009	01.0439	450154	01.1399
440087	00.9001	440167	01.1917	450029	01.1249	2009	01-1038	201	01.0603
6004	01.0622	440168	01.0226	450031	01.2116	450095	01.1121	201	01.0534
6004	01.2863	440170	00.7451	450032	01.1404	5003	01.3086	201	00.8806
600%	1996.00	440171	01.0508	450033	CI.3767	600	01.2123	501	01.2607
0104	1496.00	440173	01.0880	450034	01,3318	0	01.0363	501	01.0273
440101	00.8865	440114	00.9140	450035	01.2225	0	01.0800	450164	00.9707
4010	0.052	440175	01.0026	450037	01.2061	0	01.2188	501	0100116
104	400	440176	010.1213	450039	01.1476	450102	01.3907	450166	0100010
3 (01.2753	111044	00.9531	040040	1787-10	0100	01.0800	450109	0008746
0104	41000	821044	0101010	450041	00.4800	0100	0077010	450170	STOIS TO
0104	1 0013	407	000 4880	2400047	01.0000	0	1894-00	420114	0101313
011044		440181	00.0331	450043	01.1908	420103	01.00441	450175	01.0000
1104	+61.1	791044	1916-000	*****	0104003	430110	40130	430110	0141010

×	01.1167	01100123	1026-00	01.1233	01.0053	00.9682	01.3627	00.9293	01.3035	01.1347	0101618	0101275	01.1312	0102164	01.1168	00.9582	01.0834	00.8825	00.9455	00.8803	01-1004	00°0546	01.0851	01-0406	000000000000000000000000000000000000000	00.9256	0101711	01.0554	01-1102	01.0530	00.7884	0101319	01-1853	00.9123	00.9051	00.9774	048	00.9595	01.1858	01.0360	00.9874	01.0205
W	450544	450545	450547	450550	450551	450557	450558	450559	450561	450563	5056	450568	450570	450571	450573	450574	450575	5057	450578	450580	450581	450583	450584	450586	450587	450590	450591	450595	450596	450591	450603	450604	450605	5060	450640	450613	450614	450615	450617	450620	450621	450623
CASE MIX	01.0627	0100000	01-1490	00.9767	01.0371	01-2970	01.0640	00.9358	1766-00	01-2005	048	00.9355	00-9148	01-3066	00.9134	01.0312	00.9352	01.2967	00.9684	01-0004	01.2275	01-0075	01.0336	01.0723	01-0348	01-1506	01.0788	01.0535	01.0372	00.9440	01.0992	00.9836	01.2090	00.9629	01.1218	01.1228	01.2442	01.0648	01.2389	00.8826	010010	0102010
PROVIDER	450421	450422	450424	450425	450429	450431	450438	4	450446	450447	450450	154064	450454	450457	450458	450459	450460	450462	450464	450467	450469	450472	450473	450475	450481	450484	450486	450488	450489	450493	450497	450498	450508	450513	420014	450518	450523	450527	450530	450534	450535	450537
CASE MIX	01.0831	01.0342	01.3465	01.1451	00.9314	00.9598	01.0288	01-1683	01-1071	01.0527	01-1049	01.1183	01-1623	00-9210	01.0638	01.5212	00.8387	01.0152	01.0574	01-10-10	01.1107	01-0093	01.2794	01.1329	146000	01.2182	01.1656	01.0435	1646.00	01.1325	01-1324	01-1955	01-1565	01.0060	0000000	01-1593	01.2902	01.0241	1696.00	01-1061	01.1552	4406.00
PROVIDER	450333	450534 480227	45033B	450340	450341	450345	450343	450346	450347	450348	450349	450351	450352	450355	450357	450358	450359	450362	450365	450360	450370	450371	450372	450373	450374	450378	450379	450381	450382	450389	450391	450393	450394	450395	450400	450402	450403	450410	450411	450415	450416	450417
CASE MIX	00.8227	1460-10	00. R&7&	01-1200	9602-00	00.9565	01-1263	1028-00	01.0369	00.9700	01.0016	01.0928	01-1590	00-9907	00.9074	01.2126	01.2293	8606.00	00.9725	01.0878	01.0791	01-1101	00.9426	01.0017	01-1463	00.9478	00.9821	00.8156	2786.00	01-0774	00.8852	01-1780	01.1986	01.00.10	01.2626	00.8514	00.8568	01.2294	01-0862	00.9503	964	01.0956
PROVIDER	450250	450535	800	505	502	450261	505	45054	205	502	450270	450271	450275	450276	450278	450280	505	505	450283	500	450289	450292	502	450294	450299	503	503	50	500	450309	450311	450312	450315	450317	415054	450321	450322	450354	450325	450327	450328	450330
Wi	01.0423		0	20		.317		01.2678			.180		010,000					01.0967	01.1492	010010	01.2209	01-1983	01.0509	01.1212	01-1052	01.0089	01.0029	01.0727	6686.00	01-1024				00.9645						1900	960	00.9841
ROV	450177	450170	450181	450182	450183	450184	450185	450187	450188	450190	450191	450192	450193	450195	450196	450197	450200	450201	450203	450506	450208	450209	450210	450211	450213	450217	450218	450219	450221	450224	450229	450230	450231	450233	450234	450236	450237	450239	450241	450242	450243	420246

NOTE: CASE MIX INDEXES DO NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS.

* CASE MIX INDEXES INCLUDE CASES RECEIVED IN HOFA CENTRAL OFFICE THROUGH JUNE. 1987.

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450627	01.1533	450687	01.0474	460012	01.3898	900064	01010	490071	01.2054
450628	00.9331	450688	01.1155	460013	01.2952	490007	01.5230	490073	01.1365
450629	00.7854	450690	01.2368	460014	01.0716	490008	01.0232	490074	01.2442
450630	01.3882	450691	01.1000	460015	01.1118	600064	01.3987	490075	01-1623
450631	01.4971	450694	91.0618	450016	00.8790	490010	01.0247	490077	01-1232
450632	01.0279	450696	01.0740	460017	01.1952	490011	01.1655	490078	1626-00
50633	01.4032	450697	01-1754	460018	\$106°00	490012	01.0107	490079	01-1084
450634	01.2546	450698	0106-00	460019	00.9833	490013	01.0665	490083	00.8081
50635	01.2429	450700	01.0037	460020	01.0245	410064	01.3900	480064	01-0249
450637	01-1526	450702	01.1811	460021	01.2095	490015	01.1826	490085	01-0396
450638	01.3057	450703	01.0714	460022	1966.00	490017	01.2300	490088	01.0651
50639	01.2233	420704	01-1712	460023	01.0732	490018	01.0920	490089	01.0124
149054	00.9445	450705	00.9351	460024	00.9932	490019	01.0917	060064	01-0723
150643	01.0852	450706	01.2215	460025	00.8125	490020	01.0688	160064	01-1674
450644	01.6439	450709	01.0559	460026	1168-00	490021	01-1263	490092	01-0734
450646	01-1825	450711	01.2006	460027	00.8269	490022	011110	490093	01.1833
150647	01.4949	450712	00.8314	460029	00.9980	490023	01.1608	460064	01-1483
150648	01.0956	450713	9691010	460030	01.0441	490054	01.3835	490095	01-1589
150649	01.0128	450715	01-1985	460032	00.9403	490027	01.0202	490097	01-0574
159054	01-1926	450716	01.0957	460033	00.8732	490028	01-1519	490098	01.0779
450652	00.9548	450717	01.2452	460035	00.9546	490029	01.0980	660064	01.0841
450653	01.1398	450718	01.0174	460036	01.0142	490030	01.0681	490100	01-1847
450654	00.9811	450719	01.1149	460037	00.9482	490031	01.0393	101064	01.0719
50656	0991-10	450722	00.9777	460039	8468 00	490032	01.5795	401064	00.7229
450658	01.0112	450723	01.2002	140094	01.1737	490033	01.1115	490105	00.8738
450659	01.3397	450724	01.3226	460042	01.2698	490035	01.1278	490106	00.9539
150660	01.2897	450725	6186.00	460043	01.1862	490037	01-1033	490107	01.0289
450661	01.1089	450726	00.8945	440094	01.0906	490038	01.0677	601064	00.9959
50662	01.2031	450727	01-1351	940094	01.0473	040064	01-1580	490110	01.0663
450665	5166 00	450728	01.0355	470001	01-1157	140064	01-1134	490111	01.0588
450666	01-1267	450729	9486 00	470003	01.5192	490045	01.1161	490112	01.4023
450667	00.9518	450730	01-1937	410004	01.0970	490043	01-1346	490113	01-1093
450668	01.3812	450732	01.0512	410005	01.1563	440064	01.1011	411064	01-0435
450669	01.1814	450733	01-1349	410006	01-1331	490045	01.1442	490115	01.0582
450670	01-1067	450734	01.0722	470008	01-1245	950064	01.2461	490116	00.9716
450671	6806-00	450735	00.8922	470010	01.0792	490041	01-1112	490117	00.9855
450672	01.3003	450736	01.0361	470011	01-1616	490048	01-1626	490118	01.4172
450673	01.1102	450737	00.7176	470012	01-1508	490050	01-1313	490119	01-1671
420674	00.9918	450738	9618-00	470013	01.0416	490052	01.2903	490120	01-1863
450675	01.1401	450739	00.9542	410015	01.1581	490053	01.1514	490122	01.0774
450676	00.9983	460001	01.4395	470016	01.0600	430054	00.9614	490123	01-1023
120677	01-2209	460003	01.3251	470018	01.0725	490055	00.8783	430154	01-1416
450678	01.2488	400094	0104050	470020	00.9722	490056	00.8580	490125	01.1815
450679	01.0097	460005	01-1769	470023	01.2053	490057	01-1768	490126	01.0692
189054	01.2332	900094	01-1713	470024	01.1035	490059	01.2292	490127	00.8949
450682	01-1232	460007	01.1617	100065	00.9344	690069	00.9143	490129	00.9823
50683	01.2233	460008	01-1513	490002	00.9819	490063	01.3520	490130	01-1460
480684	8641010	600094	01.4574	490003	00.6983	990064	01.0907	100005	01-2941
450685	01-1392	460010	01.6454	*0006*	01-1643	490067	01-1150	500002	01.265
-	1	04000			7-0-0		20000	10000	

NOTE: CASE MIX INDEXES DO NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS. : CASE MIX INDEXES INCLUDE CASES RECEIVED IN HCFA CENTRAL OFFICE THROUGH JUNE., 1987.

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R CASE MIX 01.5495	0611-10	1690-10	01-1318	01.0482	01-4603	01-1939	01.4024	1696.00	01.2579	01-7125	01.5531	01.0458	01-1027	01-1113	01-1185	01.0668	01-1639	01-1528	01-1730	6461-10	1681	01.0167	01-1531	01.2328	01-1277	9660-10	01-2866	01-1867	00.9840	01-1624	01-1624	0104177	01.0592	01.4205	01-2151	01.2630	01.0798	01.2975	01-1239	01.2881	01-1266	66	53	0110110	01-1251	01.0367
PROVIDER 520037	\$20038	520039		520042	520043	520044	520045	520047	520048	520049	520051	520053	\$20054	520056	520057	520058	520059	520060	290025	520064	520054	520068	520069	520070	520071	2007	\$20075	520076	2007	520081	2008	2008	\$20084	520087	2008	2008	5000	2009	2009	5002	5002	520096	5002	860025	520100	520101
CASE MIX 01.0970	01.0112	01.0484	01-1126	-047	01.1687	•066		.986		1646.00		4496*00	01-0082	01-1704	00.9121	01.2182	01.2163	01.1351	6997-10	177100	01-1770	01.2542	01.0261	01-1161	01.0352	01.2029	01.1566	01.1821	01.0162	01.0574	01-1885	01.3484	4991-10	01-1849	01.0045	01.0715	01-1803	01-1557	01.2609	2620-10	01.3187	01.1865	01.1948	\$697·10	01-1762	_
PROVIDER 510064	290015	510066	510068	\$10070	510071	510015	\$10014	510076	510077	510080	510081	510082	510084	510085	210086	100025	520002	520003	400025	520007	\$20008	520009	520010	520011	520012	\$20013	520014	520015	910075	520018	520019	520020	520021	520025		520025	520026	520027	520028	520026	520030	520031	550035	520033	520034	520035
CASE MIX 01.2289	01-1213	054		52	.285	.088	01-1234	.954	01-0539	01.0719	.119	.982		01-0815	0626-00	01.0927	01.3688	61.0010	101010	000-00	010010	01-0354	01.0830	01-1423	01 .0956	01-1733	00.9642	01-1163	01 1300	00-9309	01.0252	9618-00	01-1468	01.1642	01.0743	01-1042		00.9782	*96*	\$110	67110	00.000	8450-10	9.	01-1524	6
PROVIDER 510001	510002	510003	510005	1000	1000	0001	1000	510011	1001	510013	1001	210015	210016	510018	610015	270015	510022	510023	*20016	510025	510027	510028	510029	510030	510031	510033	510035	510036	510038	510040	510043	510045	940015	-	510048	210050	510052	510053	md #	510055	850016	610016	210060	0001	510062	210003
CASE MIX 01.3524	01-1522	00.8034	01.0189	980	01.0781	01.1341	1966.00	01.0514	011-1106	01-1124	01.2315	01.2230	01-1566	00.9715	01.0395	01.0146	01-1674	01.02844	0102312	OD- SARO	01-0667	01.1017	01.0536	01-1055	01.0853	00.9256	00.9576	7110-10	100.00	00.9135	01.0979	01.5001	01-1852	01.1900	01.2804	01-1508	01.2317	01.2524	00.9193	0161610	01.0013	00.000	01 1753	201110	6821.10	
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NOTE: CASE MIX INDEXES DO NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS.

: CASE MIX INDEXES INCLUDE CASES RECEIVED IN HCFA CENTRAL OFFICE THROUGH JUNE. 1987.

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NOTE: CASE MIX INDEXES DO NOT INCLUDE DISCHARGES FROM PPS-EXEMPT UNITS. : CASE MIX INDEXES INCLUDE CASES RECEIVED IN HCFA CENTRAL OFFICE THROUGH JUNE., 1987.

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Urban area (constituent counties or county equivalents)	Wage index	Urban area (constituent counties or county equivalents)	Wage index	Urban area (constituent counties or county equivalents)	Wage
Abilene, TX	0.8335	De Kalb, GA		Jefferson, AL	it of
Taylor, TX	26	Douglas, GA	The state of	Saint Clair, AL	100
Aguadilla, PR	0.4624	Fayette, GA	- 1	Shelby, AL	14 21 7
Aguada, PR	2 1000	Forsyth, GA	- MICE	Walker, AL	100
Aguadilla, PR Isabella, PR	1500	Fulton, GA	2007	Bismarck, ND	0.931
Moca, PR	ALL PARTY	Gwinnett, GA		Burleigh, ND	100
Akron, OH	1.0023	Henry, GA Newton, GA	THE PARTY	Morton, ND	
Portage, OH	1.0025	Paulding, GA	1 = 115 A	Bloomington, IN	0.921
Summit, OH	THE REAL PROPERTY.	Rockdale, GA	THE REAL PROPERTY.	Monroe, IN Bloomington-Normal, IL	0.046
Albany, GA	0.7748	Spalding, GA	210	McLean, IL	0.540
Dougherty, GA	100 100 100	Walton, GA		Boise City, ID	0.982
Lee, GA	TOTAL SECOND	Atlantic City, NJ	0.9898	Ada, ID	0.00
Albany-Schenectady-Troy, NY	0.8702	Atlantic City,	The state of	Boston-Lawrence-Salem-Lowell-	The state of the s
Albany, NY	E 23	Cape May, NJ	1 1 1 3 m	Brockton, MA	1.082
Greene, NY Montgomery, NY	Service of	Augusta, GA-SC	0.8908	Essex, MA	1000
Rensselaer, NY	A PACE OF	Columbia, GA	-	Middlesex, MA	100
Saratoga, NY	1000	McDuffie, GA Richmond, GA	10000	Norfolk, MA	
Schenectady, NY	P. Carl	Aiken, SC		Plymouth, MA	10000
Albuquerque, NM	1.0188	Aurora-Elgin, IL	1 0122	Suffolk, MA	4 074
Bernalillo, MN	1.0100	Kane, IL	1.0123	Boulder-Longmont, CO	1.071
Alexandria, LA	0.8182	Kendall, IL	(T. J. D. T.)	Bradenton, FL	0.970
Rapides, LA	3000	Austin, TX	1.0409	Manatee, FL	0.0750
Allentown-Bethlehem, PA-NJ	0.9858	Hays, TX		Brazoria, TX	0.833
Warren, NJ	Subject to	Travis, TX	10000	Brazoria, TX	0.000
Carbon, PA		Williamson, TX		Bremerton, WA	0.9407
Lehigh, PA	NOS MAY	Bakersfield, CA	1.1114	Kitsap, WA	
Northampton, PA Altoona, PA	00474	Kern, CA	-	Bridgeport-Stamford-Norwalk-	Dituing 1
Blair, PA	0.9474	Baltimore, MD	1.0178	Danbury, CT	. 1.1230
Amarillo, TX	0.0326	Anne Arundel, MD Baltimore, MD		Fairfield, CT	2000
Potter, TX	0.5520	Baltimore City, MD	BH NE	Brownsville-Harlingen, TX	0.8538
Randall, TX		Carroll, MD		Cameron, TX Bryan-College Station, TX	0.007
Anaheim-Santa Ana, CA	1.2031	Harford, MD		Brazos, TX	0.9377
Orange, CA		Howard, MD		Buffalo, NY	0.9726
Anchorage, AK	1.4619	Queen Annes, MD	5300	Erie, NY	0.0120
Anchorage, AK	2000	Bangor, ME.	0.8907	Burlington, NC	0.7548
Anderson, IN	0.9175	Penobscot, ME	0 - 2 0	Alamance, NC	- TOOLS
Anderson, SC	0.7839	Baton Rouge, LA	0.8665	Burlington, VT	0.9464
Anderson, SC	0.7039	Ascension, LA East Baton Rouge, LA	1	Chittenden, VT	
Ann Arbor, MI	1.1723	Livingston, LA	HE I	Grand Isle, VT Caguas, PR	0 1001
Washtenaw, MI		West Baton Rouge, LA	To the same of	Caguas, PR	0.4001
Anniston, AL	0.7847	Battle Creek, MI	0.9670	Gurabo, PR	Miller -
Calhoun, AL	12 242	Calhoun, MI		San Lorenz, PR	A. C.
Appleton-Oshkosh-Neenah, WI	0.9792	Beaumont-Port Arthur, TX	0.9394	Aguas Buenas, PR	THE REAL PROPERTY.
Calumet, WI	New 26	Hardin, TX	Bullet 1	Cayey, PR	THE P
Outagamie, WI Winnebago, WI	and the same	Jefferson, TX	NEED!	Cidra, PR	The same
Arecibo, PR	0.4404	Orange, TX		Canton, OH	0.9195
Arecibo, PR	0.4401	Beaver County, PA	1.0368	Carroll, OH	Cassilla
Camuy, PR	9-17	Beaver, PA Bellingham, WA	4 0000	Stark, OH	700
Hatillo, PR	1000	Whatcom, WA	. 1.0823	Casper, WY	0.9842
Quebradillas, PR		Benton Harbor, MI	0.8436	Natrona, WY Cedar Rapids, IA	0.9242
Asheville, NC	0.8501	Berrien, MI	0.0430	Linn, IA	0.9242
Buncombe, NC	United to	Bergen-Passaic, NJ	1.0299	Champaign-Urbana-Rantoul, IL	0.9141
Athens, GA	0.7710	Bergen, NJ		Champaign, IL	0.0141
Clarke, GA	THE WATER	Passaic, NJ	THE LINE	Charleston, SC	0.8467
Jackson, GA	DE SECTION	Billings, MT	0.9756	Berkeley, SC	TERES A
Madison, GA Oconee, GA	THE STATE OF	Yellowstone, MT		Charleston, SC	STATE OF
Manta, GA	0.9196	Biloxi-Gulfport, MS	. 0.8012	Dorchester, SC	Relation of the last of the la
Barrow, GA	0.8190	Hancock, MS Harrison, MS	PER PER	Charleston, WV	0.9757
Butts, GA	THE REAL PROPERTY.	Binghamton, NY	0.9107	Kanawha, WV	200
Cherokee, GA		Broome, NY	0.8107	Putnam, WV Charlotte-Gastonia-Rock Hill, NC-SC	00101
Clayton, GA	THE PARTY OF	Tioga, NY	100	Cabarrus, NC	0.6424
Cobb, GA	E PETER	Birmingham, AL	0.9226	Gaston, NC	
Coweta, GA	To the same of	Blount, AL	1 10 10 10 10 10 10 10 10 10 10 10 10 10	Lincoln, NC	A COLUMN TO THE REAL PROPERTY AND ADDRESS OF THE PARTY AND ADDRESS OF T

TABLE 4a.—WAGE INDEX FOR L AREAS—Continued	JRBAN	TABLE 4a.—WAGE INDEX FOR U	RBAN	TABLE 4a.—WAGE INDEX FOR L AREAS—Continued	JRBAN
Urban area (constituent counties or county equivalents)	- Wage index	Urban area (constituent counties or county equivalents)	Wage	Urban area (consultuent counties or county equivalents)	Wage
Mecklengburg, NC		Ellis, TX	00.93	Cumberland, NC	STATE OF
Rowan, NC	10000	- Kaufman, TX	1003 -	Fayetteville-Springdale, AR	0.749
Union, NC	1000	Rockwall, TX	1	Washington, AR	Water B.
York, SC	1	Danville, VA	0.7621	Washington, AH Flint, MI	1.145
Charlottesville, VA Albermarle, VA	0.8822	Danville City, VA	CHANGE !	Genesee, MI Shiawassee, MI Florence, AL	1000
Charlottesville City, VA		Pittsylvania, VA Davenport-Rock Island-Moline, IA-IL	0.0720	Shiawassee, MI	0.705
Fluvanna, VA	100	Scott IA	0.9739		
Greene, VA	-316	Henry, IL	1 1 1		103
Chattanooga, TN-GA	A TOTAL OF	Rock Island, IL	BRUH .	Lauderdale, AL Florence, SC	0.747
Catoosa, GA Dade, GA	FR35	Dayton-Springfield, OH	1.0107	Florence, SC	1 1/2
	P HEIGHT	Clark, OH		Fort Collins-Loveland, CO	1.025
Walker, GA Hamilton, TN	1000	Greene, OH	- NEX	Larimor, CO	The state of
Marion, TN	ENVIOLE !	Miami, OH	1	Fort Lauderdale-Hollywood-Pompa-	1 240
Sequatchie, TN	1200	Montgomery, OH Daytona Beach, FL	OPEAC	no Beach, FL	1.042
Cheyenne, WY	0.8959	Volusia FI	10000	Fort Myers-Cape Coral, FL	0 808
Laramie, WY	13000	Decatur, IL	0.8966	I as FI	the arrest
Chicago, IL	1:1211	Macon, IL	DITTO SERVE OF	Fort Pierce, FL	1.005
Cook, IL	100000	Denver, CO	1.1934		
Du Page, IL	MEDI	Adams, CO	The state of	Martin, FL St. Lucie, FL Fort Smith, AR-OK	CONTRACTOR
McHenry, IL		hiapanos, co	100 X 10 1	Fort Smith, AR-OK	0.872
Chico, CA	1.1145	Denver, CO	A COLOR	Crawford, AR	
Butte, CA Cincinnati, OH-KY-IN	1 0050	Douglas, CO	The same of	Sebastian, AH	Water Land
Dearborn, IN	1.0519	Jefferson, CO Des Moines, IA	0.0004	Sequoyah, OK Fort Walton Beach, FL	0 004
Dearborn, IN Boone, KY	1000	Dallas, IA	0.5024	Okaloosa, FL	0.0211
Gampbell, KY	1	Doll, 1A	Landa U.	Fort Wayne, IN	0 9008
Kenton, KY	03:16:15	Warren, IA	1 16	Allon IN	The state of the s
Clermont, OH	Walt !	Detriot, MI	1.0911	De Kath, IN	1382
Hamilton, OH Warren, OH	RAM	Lapeer, MI	4100	Whitley, IN	The state of
Warren, OH	0.7105	Livingston, MI		Fort Worth-Arlington, TX	0.9475
Clarksville-Hopkinsville, TN-KY Christian, KY	0.7485	Macomb, MI	100	Johnson, TX	19 34
Montgomery, TN	The same of	Monroe, MI Oakland, MI	TRAPE.	Parker, TX Tarrant, TX	1
Cleveland, OH	1.0826	Saint Clair MI	Self	Fresno, CA	1.0978
Cuyahoga, OH		Wayne, MI		Fresno, CA	1.05/
Geauga, OH	14.118	Dothan, AL	0.7892	Gadsden, AL	0.8394
Lake, OH	THE REAL PROPERTY.	Dalo At	CONTROL 12	Fe 1 41	1. 5
Medina, OH		Houston, AL		Gainesville, FL	0.8902
Colorado Springs, CO	1.0047	Dubuque, IA.	0.9712	Alachua, FL	9.00
Columbia, MO	1 0279	Dubuque, IA	0.0477	Bradford, FL Galveston-Texas City, TX	4.070
Boone, MO	1.0070	Duluth, MN-WI	0.9477	Galveston, TX	1.0782
Columbia, SC	0.8450	Douglas, Wi	Contract.	Gary-Hammond, IN	1.0415
Lexington, SC	P480 # - 1	Eau Claire, WI	0.8903	Lake, IN	1
Richland, SC		Chippewa, WI		Porter, IN	THE STATE OF THE S
Columbus, GA-AL	0.7406	Eau Claire, WI		Glens Falls, NY	0.8889
Russell, AL Chattanachea GA	Contract of the last	El Paso, TX	0.8849	Warren, NY	WU.
Chattanoochee, GA Muscogee, GA	1	El Paso, TX Elkhart-Goshen, IN	0.9142	Washington, NY Grand Forks, ND	0010
Columbus, OH	0.9296	Elkhart, IN	0.9142	Grand Forks, ND.	0.9462
Delaware, OH	0.0250	Elmira, NY	0.9152	Grand Rapids, MI	1.0058
Fairfield, OH	1000	Chemung, NY	0.0102	Kent, MI	1.0000
Franklin, OH	H. Marini	Enid, OK	0.9125	Ottawa, MI	1
Licking, OH	The state of	Garfield, OK	THE REAL PROPERTY.	Great Falls, MT	0.9966
Madison, OH	200	Erie, PA	0.9488	Cascade, MT	1
Pickaway, OH	1200	Erie, Pa	+ 0000	Greeley, CO	1.0174
Union, OH Corpus Christi, TX	0.8901	Eugene-Springfield, OR	1.0353	Weld, CO	0.0000
Nueces, TX	0.000	Lane, OR Evansville, IN-KY	0.9963	Green Bay, WI	-
San Patricio, TX	TOTAL T	Posev IN		Greensboro-Winston-Salem-High	DESCRIPTION OF
Cumberland, MD-WV	0.8798	Vanderburgh, IN	SE SE	Point, NG	0.8710
Allegeny, MD	MENTER:	Marrick IN		Davidson, NC	1 11213
Mineral, WV		Henderson, KY		Davie, NC	
Dallas, TX	0.9565	Fargo-Moorhead, ND-MN	1 0031	Forsyth, NC	estreets.
Collin, TX Dallas, TX	PERSONAL PROPERTY.	Clay, MN	THE PLANT	Guillord, NC	100
	SCHOOL S	Cass, ND Fayetteville, NC	Contrary .	Randolph, NC Stokes, NC	

110	_	The state of the s		AREAS—Continued	-
Urban area (constituent counties or county equivalents)	Wage	Urban area (constituent counties or county equivalents)	Wage	Urban area (constituent counties or county equivalents)	Wag
Yadkin, NC	The same of	Onslow, NC	THE PERSON	Lancaster, PA	Date:
Greenville-Spartanburg, SC	0.8961	Janesville-Beloit, WI	0.8935	Lansing-East Lansing, MI	1.025
Greenville, SC	HERE W	Rock, WI	THE PERSON NAMED IN	Clinton, MI	
Pickens, SC	4 7 2 2	Jersey City, NJ	1.0599	Eaton, MI	-
Spartanburg, SC	0.0000	Hudson, NJ	10011	Ingham, MI	
Hagerstown, MDWashington, MD	0.8869	Johnson City-Kingsport-Bristol, TN-	2 40000	Laredo, TX	0.75
Hamilton-Middletown, OH	0.0640	VA	0.8446	Webb, TX	300
Butler, OH	0.9049	Hawkins, TN	100	Las Cruces, NM	0.83
Harrisburg-Lebanon-Carlisle, PA	0.9907	Sullivan, TN	400 4 1	Dona Ana, NM Las Vegas, NV	4 000
Cumberland, PA	and the same of th	Unicoi, TN	SE TON	Clark, NV	1.08
Dauphin, PA	THE REAL PROPERTY.	Washington, TN	Bled t	Lawrence, KS	0.97
Lebanon, PA		Bristol City, VA	5000	Douglas, KS	0.97
Perry, PA	19815	Scott, VA		Lawton, OK	0.85
lartford-Middletown-New Britain-	A STATE OF THE	Washington, VA	100 MIN TO 1	Comanche, OK	0.00
Bristol, CT	1.0898	Johnstown, PA	0.9060	Lewiston-Auburn, ME	0.90
Hartford, CT	COLD ST	Cambria, PA	TO THE	Androscoggin, ME	0.00
Litchfield, CT		Somerset, PA	PER SE	Lexington-Fayette, KY	0.92
Middlesex, CT		Joliet, IL	1.0507	Bourbon, KY	100000000000000000000000000000000000000
Tolland, CT		Grundy, IL	TEST WILLIAM	Clark, KY	1
Hickory, NC	0.8335	Will, IL	All the second	Fayette, KY	To Take
Alexander, NC Burke, NC		Joplin, MO	0.8649	Jessamine, KY	1
Catawba, NC		Jasper, MO	The state of the s	Scott, KY	
Honolulu, HI	1 1240	Newton, MO		Woodford, KY	Transaction and
Honolulu, HI	1.1343	Kalamazoo, MI	1.1352	Lima, OH.	0.923
louma-Thibodaux, LA	0.8083	Kalamazoo, MI Kankakee, IL	0.0000	Allen, OH	
Lafourche, LA	0.0003	Vankakaa II	To be a second	Auglaize, OH	
Terrebonne, LA		Kansas City, KS-MO	1 0064	Lincoln, NE	0.928
louston, TX	0.9868	Johnson, KS	1.0004	Lancaster, NE Little Rock-North Little Rock, AR	0.000
Fort Bend, TX	CONTRACTOR OF	Leavenworth, KS		Faulkner, AR	0.937
Harris, TX		Miami, KS	dina:	Lonoke, AR	- DRIV
Liberty, TX		Wyandotte, KS		Pulaski, AR	10000
Montgomery, TX		Cass, MO		Saline, AR	THE !
Waller, TX	No. of the last	Clay, MO		Longview-Marshall, TX	0.803
funtington-Ashland, WV-KY-OH	0.9066	Jackson, MO	A THE STREET	Gregg, TX	
Boyd, KY	15000	Lafayette, MO		Harrison, TX	1
Carter, KY	SAME AND A	Platte, MO		Lorain-Elyria, OH	0.951
Greenup, KY	DENEST'S.	Ray, MO		Lorain, OH	133
Lawrence, OH Cabell, WV	13800	Kenosha, WI	1.0384	Los Angeles-Long Beach, CA	1.246
Wayne, WV		Kenosha, WI		Los Angeles, CA	
funtsville, AL	0.8208	Killeen-Temple, TX	0.9789	Louisville, KY-IN	0.952
Madison, AL	0.0200	Bell, TX Coryell, TX		Clark, IN	1
ndianapolis, IN	0.9941	Knoxville, TN	0.0005	Floyd, IN	-innie
Boone, IN	0.0041	Anderson, TN	0.8335	Harrison, IN	and the
Hamilton, IN		Blount, TN	The state of the	Bullitt, KY Jefferson, KY	WHO IS
Hancock, IN	A457 1	Grainger, TN	Colonia in	Oldham, KY	0000
Hendricks, IN	bristone,	Jefferson, TN	No a	Shelby, KY	Which's
Johnson, IN	idmir)	Knox, TN		Lubbock, TX	0.956
Marion, IN	SINGE !	Sevier, TN	Brus :	Lubbock, TX	0.000
Morgan, IN	1 11977	Union, TN	HIESON.	Lynchburg, VA	0.858
Shelby, IN		Kokomo, IN	0.9352	Amherst, VA	
owa City, IA	1.1630	Howard, IN	Lines I.	Campbell, VA	Samuel Samuel
Johnson, IA		Tipton, IN	A COLUMN TO	Lynchburg City, VA	To Bu
ackson, MI	0.9445	LaCrosse, WI	0.9629	Macon-Warner Robins, GA	0.828
ackson, MS	0.0400	LaCrosse, WI		Bibb, GA	1 130
Hinds, MS	0.8439	Lafayette, LA	0.9261	Houston, GA	
Madison, MS		St. Martin, LA	Tanta E	Jones, GA	
Rankin, MS	Name of	Lafayette, IN	0.9796	Peach, GA	4.040
ackson, TN	0.7506	Tippecanoe, IN	0.8736	Madison, WI	1.016
Madison, TN	0.7000	Lake Charles, LA	0.0170	Dane, WI	0.000
acksonville, FL	0.8923	Calcasieu, LA	0.9172	Manchester-Nashua, NH	0.922
Clay, FL	1	Lake County, IL	1.0904	Hillsborough, HN Merrimack, NH	Stone .
Duval, FL	10000	Lake, IL	1.0004	Mansfield, OH	0.911
Nassau, FL	News of	Lakeland-Winter Haven, FL	0.8261	Richland, OH	0.911
St. Johns, FL	1 200	Polk, FL		Mayaguez, PR	0.484
acksonville, NC	0.7358	Lancaster, PA	0.0888	Anasco, PR	3,104

TABLE	4a1	NAGE	NDEX	FOR	URBAN
	ARE	AS-C	ontinu	led	

There of third of	
Urban area (constituent counties or county equivalents)	Wage index
Cabo Rojo, PR	The said
Hermigueros, PR	Dian.
Mayaguez, PR	Om O
San German, PR	
McAllen-Edinburg-Mission, TX Hidalgo, TX	0.7655
Medford, OR	0.9701
Jackson, OR	and the bare
Melbourne-Titusville, FL	0.8862
Brevard, FL Memphis, TN-AR-MS	0.9644
Crittenden, AR	0.9044
De Soto, MS	Section 1
Shelby, TN	Technical v
Tipton, TN	4.0707
Merced, CA	1.0727
Merced, CA Miami-Hialeah, FL	1.0151
Dade, FL	
Middlesex-Somerset-Hunterdon, NJ	0.9837
Hunterdon, NJ Middlesex, NJ	THE PERSON
Somerset, NJ	Service of
	1.0578
Midland TX	
Milwaukee, WI	1.0435
Ozaukee, WI	SPINIS C
Washington, WI	SHOULD
Waukesha, WI	DEPOSED T
Minneapolis-St. Paul, MN-WI	1.1224
Carver, MN	DIOVICE IV
Chisago, MN	
Dakota, MN	(Indignal)
Hennepin, MN Isanti, MN	3000
Ramsey, MN	E COLOR
Scott, MN	
Washington, MN Wright, MN	CARACI
St. Croix, WI	Market 1
	0.8319
Baldwin, AL	Topid
Mobile, AL Modesto, CA	1.1049
Stanislaus, CA	1.1049
Monmouth-Ocean, NJ	0.9365
Monmouth, NJ	10-10
Ocean, NJ Monroe, LA	0.8471
Quachita I.A	V. O. T.
Montgomery, AL	0.8173
Autauga, AL Elmore, AL	dimini -
Elmore, AL Montgomery, AL	All Division in
Muncie, IN	0.9565
Delaware, IN	0.0000
Muskegon, MI Muskegon, MI	The same of the sa
Naples, FL	0.9919
Collier, FL	C. month
Nashville, TN	Annual Contract of the Contrac
Cheatham, TN Davidson, TN	
Dickson, TN	English
Robertson, TN Rutherford, TN	201201
Rutherford, TN Sumner, TN	THE PARTY
The state of the s	STATE OF THE PARTY

TABLE 4a.—WAGE INDEX FOR URBAN AREAS—Continued

AREAS—Continued					
Urban area (constituent counties or county equivalents)	Wage				
Williamson, TN Wilson, TN					
Wilson, TN Nassau-Suffolk, NY Nassau, NY	1.2359				
Suffolk, NY New Bedford-Fall River-Attleboro,	Total Control				
MA	0.9352				
New Haven-Waterbury-Meriden, CT New Haven, CT	CALCOL .				
New London-Norwich, CT New London, CT	Made 1				
New Orleans, LA Jefferson, LA Orleans, LA	0.9080				
	elegion				
St. John The Baptist, LA St. Tammany, LA					
New York, NY	1.3092				
Kings, NY New York City, NY	Al pelgoli Manual				
Queens, NY	Obert .				
Minetahantar MV					
Newark, NJ Essex, NJ	and the second				
Morris, NJ Sussex, NJ	THE REAL PROPERTY.				
	0.8492				
Niagara, NY Norfolk-Virginia Beach Newport News, VA					
Chesapeake City, VA Gloucester, VA	0.9196				
Hampton City, VA James City Co., VA	artomial.				
Newport News City, VA Norfolk City, VA	Caracter N				
Poquoson, VA Portsmouth City, VA	DE STATE OF THE ST				
Suffolk City, VA Virginia Beach City, VA Williamsburg City, VA	anticus .				
York, VA	1,4023				
Alameda, CA Contra Costa, CA	aparti.				
Ocala, FL Marion, FL Odessa, TX					
	TANK BE				
Canadian, OK Cleveland, OK					
McClain, OK					
Oklahoma, OK Pottawatomie, OK Olympia, WA	PAGE NELL				
Thurston, WA Omaha, NE-IA	0.9822				
Pottawattamie, IA Douglas, NE	Today				
Sarpy, NE	DESIGNATION OF				

TABLE 4a.—WAGE INDEX FOR URBAN AREAS—Continued

	-
Urban area (constituent counties or county equivalents)	Wage
County Coparation (Control	
Washington, NE	
Orange County, NY	0.8828
Orange, NY Orlando, FL	0.9356
Orange FI	
Oscania FI	
Seminole, FL Owensboro, KY	0.0000
Owensboro, KY Daviess, KY	0.8360
Oxnard-Ventura, CA	1.2976
Ventura CA	
Panama City, FL	0.7882
Bay, FL Parkersburg-Marietta, WV-OH	0.8828
Washington, OH	GIOULS
Wood, WV	100000
Pascugoula, MS	0.8929
Jackson, MS Pensacola, FL	0.8241
Fecambia FI	N ROLL
Santa Rosa, FL	CHESTON
Peoria, IL	0.9879
Tazewell, IL	Calend
Woodford, IL	DESIGNATION AND ADDRESS OF THE PERSON ADDRESS OF THE PERSON AND ADDRESS OF THE PERSON ADDR
Philadelphia, PA-NJ	1.0935
Burlington, NJ Camden, NJ	Muole I
Gloucester, NJ	daner -
Bucks, PA	
Chester, PA	
Delaware, PA Montgomery, PA	
Philadelphia, PA	Tanada .
Phoenix AZ	1.0079
Maricopa, AZ	0.7707
Pine Bluff, AR	
Pittsburgh, PA	1.0240
Allegheny, PA	Alson I
Fayette, PA	
Washington, PA Westmoreland, PA	TO HAVE
Pittsfield, MA	0.9946
Berkshire, MA	
Ponce, PRJuana Diaz, PR	0.5513
Ponce PR	MINE S
Portland, ME	0.9461
Cumberland, ME	THE PERSON NAMED IN
Sagadahoc, ME York, ME	SUM!
Portland, OR	1 1292
Clackamas, OR	III OF STREET
Multnomah, OR Washington, OR	-29
Yambill OR	DOMESTIC:
Portsmouth-Dover-Rochester, NH	0.9114
Rockingham, NH	THE PARTY
Strafford, NH Poughkeepsie, NY	0.9597
Dutchess, NY	0.3331
Providence-Pawtucket-Woonsocket,	eshowy
RI	0.9811
Bristol, RI Kent, RI	THE PARTY OF
Newport, RI	Sans.
Providence, RI	WUE.
Washington, RI	Deliber

TABLE 4a.—WAGE INDEX FOR AREAS—Continued	OHBAN	TABLE 4a.—Wage Index FOR I	JRBAN	TABLE 4a.—Wage Index For I	URBAN
Urban area (constituent counties or county equivalents)	Wage index	Urban area (constituent counties or county equivalents)	Wage index	Urban area (constituent counties or county equivalents)	Wage
Provo-Orem, UT	0.9278	Buchanan, MO		S C4	POR
Utah, UT	A SECTION	St. Louis, MO-IL	1 0165	Sonoma, CA Sarasota, FL	0.016
Pueblo, CO	0.9920	Clinton, IL	1.0105	Sarasota, FL	0.916
Pueblo, CO	the property	Jersey, IL	Time II	Savannah, GA	0.840
Racine, WI	0.9299	Madison, IL	Louis .	Chatham, GA	0.040
Racine, WI	PERS.	Monroe, IL	10000	Effingham, GA	Town.
Raleigh-Durham, NC Durham, NC	0.9274	St. Clair, IL	pinze -	Scranton-Wilkes Barre, PA	0.931
Franklin, NC	The second second	Franklin, MO	1-10/2019	Columbia, PA	America .
Orange, NC	Tana hal	Jefferson, MO St. Charles, MO	1000	Lackawanna, PA	man of
Wake, NC	-	St. Louis, MO	THE STATE OF THE S	Luzerne, PA	DONN.
Rapid City, SD	0.8702	St. Louis City, MO	-	Monroe, PA Wyoming, PA	- HONOR
Pennington, SD	Mouthwelle.	Salem, OR	1 0416	Seattle, WA	1 000
Reading, PA	0.9381	Marion, OR	1.0410	King, WA	1.090
Berks, PA	Estimate	Polk, OR	2000	Snohomish, WA	1
Redding, CA	1.0779	Salinas-Seaside-Monterey, CA	1.2211	Sharon, PA	0.919
Shasta, CA		Monterey, CA	AND ROOM	Mercer, PA	0.010
Reno, NV	1.1202	Salt Lake City-Ogden, UT	0.9508	Sheboygan, WI	0.931
Washoe, NV	A COUNTY	Davis, UT	andstyl	Sheboygan, WI	1
Richland-Kennewick, WA	0.9688	Salt Lake, UT	0207	Sherman-Denison, TX	0.828
Benton, WA Franklin, WA	1000	Weber, UT		Grayson, TX	M. mari
Richmond-Petersburg, VA	0.0007	San Angelo, TX	0.8302	Shreveport, LA	0.899
Charles City Co., VA	0.8897	Tom Green, TX	Ignini V	Bossier, LA	P. Bear
Chesterfield, VA	10000	San Antonio, TX	0.8377	Caddo, LA	POLICIES.
Colonial Heights City, VA		Bexar, TX Comal, TX	The state of the s	Sioux City, IA-NE	0.924
Dinwiddie, VA		Guadalupe, TX	THE PARTY	Woodbury, IA	1-26-01
Goochland, VA	1	San Diego, CA	1 2250	Dakota, NE Sioux Falls, SD	0.055
Hanover, VA	A COUNTY	San Diego, CA	1.2000	Minnehaha,SD	0.9552
Henrico, VA	TE CHINA	San Francisco, CA	1.4946	South Bend-Mishawaka, IN	0.960
Hopewell City, VA	1000	Marin, CA	300	St. Joseph, IN	0.300
New Kent, VA	BEALES .	San Francisco, CA	WAR I	Spokane, WA	1.0823
Petersburg City, VA	resond.	San Mateo, CA	AU SEN I	Spokane, WA	1.00
Powhatan, VA	F-500-5	San Jose, CA	1.4323	Springfield, IL	1.0040
Prince George, VA Richmond City, VA	Punton	Santa Clara, CA	DING 1	Menard, IL	- District
Riverside-San Bernardino, CA	1 1500	San Juan, PR	0.5387	Sangamon, IL	DOM:
Riverside, CA	1.1000	Barcelona, PR Bayoman, PR	DIENES I	Springfield, MO	0.9074
San Bernardino, CA	Halipa	Canovanas, PR	DOMESTIC OF	Christian, MO Greene, MO	To be A
Roanoke, VA	0.8346	Carolina, PR	THE REAL PROPERTY.	Springfield, MA	0.0350
Botetourt, VA	200000	Catano, PR	20050	Hampden, MA	0.8750
Roanoke, VA	The same of the	Corozal, PR		Hampshire, MA	Service Co.
Roanoke City, VA	Land Color	Dorado, PR	111123	State College, PA	1.0303
Salem City, VA		Fajardo, PR	WEER	Centre, PA	PERCHAIL.
Rochester, MNOlmsted, MN	1.0027	Florida, PR		Steubenville-Weirton, OH-WV	0.9106
Rochester, NY	0.0550	Guaynabo, PR	733	Jefferson, OH	Mount
Livingston, NY	0.9558	Humacao, PR	H (18)	Brooke, WV	10000
Monroe, NY	154 2145	Juncos, PR		Hancock, WV	100000
Ontario, NY		Los Piedras, PR Loiza, PR		Stockton, CA	1.1743
Orleans, NY		Luguillo, PR		San Joaquin, CA	0.070
Wayne, NY	65	Manati, PR		Syracuse, NY	0.9730
Rockford, IL	1.0245	Naranjito, PR	THE REAL PROPERTY.	Onondaga, NY	E PROPERTY OF THE PARTY OF THE
Boone, IL	The state of the s	Rio Grande, PR	- manny	Oswego, NY	STREET, SQUARE,
Winnebago, IL	1	San Juan, PR		Tacoma, WA	1.0325
Sacramento, CA	1.2140	Toa Alta, PR		Pierce, WA	1.0020
Eldorado, CA		Toa Baja, PR	MEET	Tallahassee, FL	0.8531
Placer, CA	100 mm	Trojillo Alto, PR	E REEL	Gadsden, FL	
Sacramento, CA	1 3 1 1 1	Vega Alta, PR	STATE	Leon, FL	100
Yolo, CA Saginaw-Bay City-Midland, MI	1.0507	Vega Baja, PR	11314	Tampa-St. Petersburg-Clearwater,	The state of the s
Bay, MI	1.0597	Santa Barbara-Santa Maria-Lompoc,	N. Alessain	FL	0.9125
Midland, MI	Here is	CA	1.1428	Hernando, FL	
Saginaw, MI		Santa Barbara, CA Santa Cruz, CA	1.2017	Hillsborough, FL	1000
St. Cloud, MN	0.9662	Santa Cruz, CA	1.2017	Pasco, FL Pinellas, FL	
Benton, MN		Santa Fe, NM	0.9362	Terre Haute, IN	0.8090
Sherburne, MN	The State of	Los Alamos, NM	5.505E	Clay, IN	0.8090
Stearns, MN	100000	Santa Fe, NM	9012.0	Vigo, IN	TEN -
St. Joseph, MO	0.8811	Santa Rosa-Petaluma, CA	1 2043	Texarkana-TX-Texarkana, AR	0.0074

TABLE 4a.—WAGE INDEX FOR URBAN AREAS—Continued

Urban area (constituent counties or county equivalents)	Wage
Miller, AR	162
Bowie, TX	JES M.
Toledo, OH	1.1101
Fulton, OH	WILL THE STREET
Lucas, OH	12000
Wood, OH	CALL .
Topeka, KS	0.9955
Shawnee, KS	HINE A
Trenton, NJ	1.0014
Mercer, NJ	
Tucson, AZ	0.9639
Pima, AZ	0.0040
Tulsa, OK	0.9346
Creeks, OK	Town St.
Osage, OK Rogers, OK	PER COLUMN
Tulsa, OK	Sec.
Wagoner, OK	10000
Tuscaloosa, AL	0.9515
Tuscaloosa, AL	0.0010
Tyler, TX	0.9326
Smith, TX	0,000
Utica-Rome, NY	0.8211
Herkimer, NY	AL VI
Oneida, NY	4300F
Vallejo-Fairfield-Napa, CA	1.2767
Napa, CA	Bull -
Solano, CA	111111111111111111111111111111111111111
Vancouver, WA	1.0772
Clark, WA	-
Victoria, TX	0.7993
Victoria, TX	0.0000
Vineland-Millville-Bridgeton, NJ	0.9580
Cumberland, NJ Visalia-Tulare-Porterville, CA	4 4 4 4 4 0
Tulare, CA	1.1418
Waco, TX	0.8585
McLennan, TX	0.0000
Washington, D.CMD-VA	1.1051
District of Columbia, DC	1.103
Calvert, MD	ORT IT
Charles, MD	OTOT
Frederick, MD	C. HOL
Montgomery, MD	Many .
Prince Georges, MD	minutes and
Alexandria City, VA	Blatter A
Arlington, VA	THE REAL PROPERTY.
Fairfax, VA	The second
Fairfax City, VA	Di -chi
Falls Church City, VA	THE L
Loudoun, VA	13373
Manassas City, VA	2001
Manassas Park City, VA	1

TABLE 4a.—WAGE INDEX FOR URBAN AREAS—Continued

Urban area (constituent counties or county equivalents)	Wage
Prince William, VA	
Stafford, VA	
Waterloo-Cedar Falls, IA	0.9432
Black Hawk, IA	0.0102
Bremer, IA	
Wausau, WI	0.9457
Marathan, WI	20.70
West Palm Beach-Boca Raton-	
Delray Beach, FL	0.9431
Palm Beach, FL	
Wheeling, WV-OH	0.8761
Belmont, OH	
Marshall, WV	
Ohio, WV	
Wichita, KS	1.0469
Butler, KS	
Harvey, KS	
Sedgwick, KS	
Wichita Falls, TX	0.8221
Wichita, TX	
Williamsport, PA	0.8804
Lycoming, PA	
Wilmington, DE-NJ-MD	1.0125
New Castle, DE	
Cecil, MD	
Salem, NJ	
Wilmington, NC	0.8602
New Hanover, NC	
Worcester-Fitchburg-Leominster, MA	0.9460
Worcester, MA	
Yakima, WA	0.9850
Yakima, WA	
York, PA	0.9340
Adams, PA	
York, PA	The same
The state of the s	0.9942
Mahoning, OH	
Trumball, OH	PRINCES IN
Yuba City, CA	0.9970
Sutter, CA	
Yuba, CA	
Control of the last of the las	

TABLE 4b—WAGE INDEX FOR RURAL AREAS

Nonurban area	Wage index
Alabama	0.7005
Alaska	1.3922
Arizona	0.8869
Arkansas	0.7124

TABLE 4b—WAGE INDEX FOR RURAL AREAS—Continued

Nonurban area	Wage index
California	1.0400
California	1.0428
Colorado	0.8666
Connecticut	1.0013
Delaware	0.8236
Florida	0.8223
Georgia	0.7385
Hawaii	0.9318
Idaho	0.8489
Illinois	0.8188
Indiana	0.8104
lowa	0.8070
Kansas	0.7927
Kentucky	0.7754
Louisiana	0.7856
Maine	0.8191
Maryland	0.8112
Massachusetts	1.0033
Michigan	0.9036
Minnesota	0.8605
Mississippi	0.7215
Missouri	0.7640
Montana	0.8558
Nebraska	0.7751
Nevada	0.9817
New Hampshire	0.8784
New Jersey 1	
New Mexico	0.8359
New York	0.8124
North Carolina	0.7650
North Dakota	0.8463
Ohio	0.8609
Oklahoma	0.7938
Oregon	1.0029
Pennsylvania	0.8807
Puerto Rico	0.5536
Rhode Island 1	
South Carolina	0.7232
South Dakota	0.7668
Tennessee	0.7162
Texas	0.7591
Utah	0.8782
Vermont	0.8387
Virginia	0.7833
Virgin Islands 1	
Washington	0.9806
West Virginia	0.8414
Wisconsin	0.8458
Wyoming	0.9100

¹ All counties within the State are classified urban.

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	STAY DUILLIER CUTDER POINTS USED IN THE PROSPECTIVE PAYMENT SYSTEM
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ARITHHETIC NEAN LOS 18.8 18.3 16.9 8.0	20.3 20.3 5.1 11.1	7.99.7 9.90.0 9.90.0	9.0 6.0 11.2	25.5	3.6 88.9 5.2	6.0 3.7 5.6
RELATIVE MEIGHTS 3.4434 3.8160 2.9183 2.5904 1.5685	2.5269 2.5269 7367 1.2639 1.2123	.9459 .9324 1.2429	1.0384 .6358 .9557 .6158	1.3613 .7055 .9505 .9228 .5386	1.4753 1.1694 .5856 .3539	.6550 .4005 .2457 1.2038
AND HOLDS WANGER TO AND	MITH CC	S	A GREAT TOP DATE		3 9	
G CRANIDTONY AGE >17 EXCEPT FOR TRAUMA. G CRANIDTOMY FOR TRAUMA AGE >17 G * CRANIDTOMY AGE 0-17 G SPINAL PROCEDURES G EXTRACRANIAL VASCULAR PROCEDURES	CARPAL TUNNEL RELEASE PERIPH & CRANIAL NERVE & OTHER NERV SYST PROC PERIPH & CRANIAL NERVE & OTHER NERV SYST PROC SPINAL DISORDERS & INJURIES NERVOUS SYSTEM NEOPLASHS MITH CC	NERVOUS SYSTEM NEOPLASMS M/O CC DEGENERATIVE NERVOUS SYSTEM DISORDERS MULTIPLE SCLEROSIS & CEREBELLAR ATAXIA SPECIFIC CEREBROVASCULAR DISORDERS EXCEPT TIA TRANSIENT ISCHEMIC ATTACK & PRECEREBRAL OCCLUSIONS	NONSPECIFIC CERFBROVASCULAR DISCROBERS W CC NONSPECIFIC CEREBROVASCULAR DISCROBERS W/O CC CRANIAL & PERIPHERAL NERVE DISCROBERS WITH CC CRANIAL & PERIPHERAL NERVE DISCROBERS W/O CC NERVOUS SYSTEM INFECTION EXCEPT VIRAL MENINGITIS	VIRAL MENINGITIS HYPERTENSIVE ENCEPHALOPATHY NONTRAUMATIC STUPOR & COMA SEIZURE & HEADACHE AGE >17 MITH CC SEIZURE & HEADACHE AGE >17 M/O CC	SEIZURE & HEADACHE AGE 0-17 IRAUMATIC STUPOR & COMA, COMA <1 HR AGE >17 MITH CC IRAUMATIC STUPOR & COMA, COMA <1 HR AGE >17 MITH CC TRAUMATIC STUPOR & COMA, COMA <1 HR AGE >17 M/O CC * TRAUMATIC STUPOR & COMA, COMA <1 HR AGE >17	CONCUSSION AGE >17 WITH CC CONCUSSION AGE >17 W/O CC + CONCUSSION AGE 0-17 OTHER DISORDERS OF NERVOUS SYSTEM WITH CC OTHER DISORDERS OF NERVOUS SYSTEM W/O CC
SURG CRANIDTONY AGE >17 EXCEPT FOR SURG CRANIDTONY AGE >17 SURG * CRANIDTONY AGE 0-17 SURG SPINAL PROCEDURES SURG EXTRACRANIAL VASCULAR PROCEDUR	SURG CARPAL TUNNEL RELEASE SURG PERIPH & CRANIAL NERVE & OTHER NERV SYST PROC SURG PERIPH & CRANIAL NERVE & OTHER NERV SYST PROC MED SPINAL DISORDERS & INJURIES MED NERVOUS SYSTEM NEOPLASHS MITH CC	MED VERYDUS SYSTEM NEDPLASMS M/C MED DEGENERATIVE NERVOUS SYSTEM MED MULTIPLE SCLEROSIS & CEREBEL MED SPECIFIC CEREBROYASCULAR DIS MED TRANSIENT ISCHEMIC ATTACK &	MED NUNSPECIFIC CEREBROVASCULAR DISCROBERS MED NUNSPECIFIC CEREBROVASCULAR DISCROBERS MED CRANIAL & PERIPHERAL NERVE DISCROBERS MED CRANIAL & PERIPHERAL NERVE DISCROBERS MED NERVOUS SYSTEM INFECTION EXCEPT VIRAL	MED	MED SEIZURE & HEADACHE AGE 0-17 MED TRAUMATIC STUPOR & COMA, COMA >1 HR AGE MED TRAUMATIC STUPOR & COMA, COMA <1 HR AGE MED * TRAUMATIC STUPOR & COMA, COMA <1 HR AGE MED * TRAUMATIC STUPOR & COMA, COMA <1 HR AGE	CONCUSSION AGE >17 MITH CC CONCUSSION AGE >17 M/O CC CONCUSSION AGE O-17 OTHER DISORDERS OF NERVOUS SYSTEM OTHER DISORDERS OF NERVOUS SYSTEM
CRANIDTONY AGE >17 EXCEPT FOR CRANIDTOMY FOR TRAUMA AGE >17 SPINAL PROCEDURES EXTRACRANIAL VASCULAR PROCEDUR	CARPAL TUNNEL RELEASE PERIPH & CRANIAL NERVE & OTHER NERV SYST PROC PERIPH & CRANIAL NERVE & OTHER NERV SYST PROC SPINAL DISORDERS & INJURIES NERVOUS SYSTEM NEOPLASHS MITH CC	NERVOUS SYSTEM NEDPLASMS W/C DEGENERATIVE NERVOUS SYSTEM MULTIPLE SCLEROSIS & CEREBEL SPECIFIC CEREBROVASCULAR DIS TRANSIENT ISCHEMIC ATTACK &	NONSPECIFIC CEREBROVASCULAR DISORDERS NONSPECIFIC CEREBROVASCULAR DISORDERS CRANIAL & PERIPHERAL NERVE DISORDERS H CRANIAL & PERIPHERAL NERVE DISORDERS M NERVOUS SYSTEM INFECTION EXCEPT VIRAL		SEIZURE & HEADACHE AGE D-17 IRAUMATIC STUPOR & COMA, COMA >1 HR AGE IRAUMATIC STUPOR & COMA, COMA <1 HR AGE ** TRAUMATIC STUPOR & COMA, COMA <1 HR AGE ** TRAUMATIC STUPOR & COMA, COMA <1 HR AGE	CONCUSSION AGE >17 WITH CC CONCUSSION AGE >17 W/O CC + CONCUSSION AGE 0-17 OTHER DISORDERS OF NERVOUS SYSTEM OTHER DISORDERS OF NERVOUS SYSTEM

* MEDICARE DATA HAVE BEEN SUPPLEMENTED BY DATA FROM MARYLAND AND MICHIGAN FOR LOW VOLUME DRGS. ** DRGS 469 AND 470 CONTAIN CASES WHICH COULD NOT BE ASSIGNED TO VALID DRGS. NOTE: GEOMETRIC MEAN IS USED DNLY TO DETERMINE PAYMENT FOR OUTLIER AND TRANSFER CASES. NOTE: RELATIVE WEIGHTS ARE BASED ON MEDICARE PATIENT DATA AND MAY NOT BE APPROPRIATE FOR OTHER PATIENTS.

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DIAGNOSIS RELATED GROUPS (DRGS), RELATIVE WEIGHTING FACTORS, ARITHMETIC AND GEOMETRIC MEAN LENGTH OF STAY, AND LENGTH OF STAY OUTLIER CUTOFF POINTS USED IN THE PROSPECTIVE PAYMENT SYSTEM 40 LIST

TIC GEDMETRIC DUTLIER LOS MEAN LOS THRESHOLD 3.6 3.1 10 4.1 3.0 14 2.8 2.2 9 2.1 1.8 5 2.6 2.0 7	3.1 2.5 4.5 3.7 14 6.8 5.6 20 4.3 1.3 14	5.6 3.8 22 3.6 2.6 13 15.7 11.9 30 3.4 2.8 9	2.9 2.3 8 3.5 2.7 11 3.1 2.3 10 2.3 1.6 6	2.4 1.8 5.1 3.2 20 2.0 1.7 5	3.2 2.1 11 7.4 4.6 23 8.5 4.7 23 4.2 3.5 12	3.9 5.8 6.1 6.1 6.0 9.2 1.3 1.3
RELATIVE ARITHMETIC MEIGHTS MEAN LOS - 6820 3.6 - 7104 4.1 - 3779 2.8 - 5167 2.1 - 4675 2.5	.3657 .6600 .3727 .6352	.6195 .3611 .4018 2.8923 .6681	. 5424 . 6159 . 6889 . 6598	.4471 .7907 .3097 .3845	.5401 .3089 1.1538 1.0548 84600	. 4272 . 9964 . 7217 . 5366 . 5345
SURG RETINAL PROCEDURES SURG ORBITAL PROCEDURES SURG PRIMARY IRIS PROCEDURES SURG LENS PROCEDURES WITH OR WITHOUT VITRECTOMY SURG EXTRADGULAR PROCEDURES EXCEPT ORBIT AGE >17	SURG * EXTRADCULAR PROCEDURES EXCEPT ORBIT AGE 0-17 SURG INTRADCULAR PROCEDURES EXCEPT RETINA, IRIS & LENS HFD HYPHEMA HED ACUTE MAJOR EYE INFECTIONS HED NEUROLOGICAL EYF DISORDERS	MED JIHER DISGROERS OF THE EYE AGE >17 M CC MED OTHER DISGROERS OF THE EYE AGE >17 M/O CC MED * OTHER DISGROERS OF THE EYE AGE 0-17 SURG MAJOR HEAD & NECK PROCEDURES SURG SIALUADENECTOMY	SURG SALIVARY GLAND PROCEDURES EXCEPT SIALGADENECTOMY SURG CLEFT LIP & PALATE REPAIR SURG SINUS & MASTOID PROCEDURES AGE >17 SURG * SINUS & MASTOID PROCEDURES AGE 0-17 SURG * MISCELLANEOUS EAR, NOSE & THROAT PROCEDURES	SURG TEA PROC. EXCEPT TONSILLECTOMY E/OR ADENDIDECTOMY ONLY. AGE >17 SURG * TEA PROC. EXCEPT TONSILLECTOMY E/OR ADENDIDECTOMY ONLY. AGE 0-17 SURG * TONSILLECTOMY E/OR ADENDIDECTOMY ONLY. AGE >17 SURG * TONSILLECTOMY E/OR ADENDIDECTOMY ONLY. AGE 0-17	SURG * MYRINGOTOMY M TUBE INSERTION AGE >17 SURG * MYRINGOTOMY M TUBE INSERTION AGE 0-17 SURG OTHER EAR. NOSE & THROAT O.R. PROCEDURES MED EAR. NOSE & THROAT MALIGNANCY MED DYSEQUILIBRIUM	ED EPISTAXIS CO CITIES HEDIA & URI AGE >17 WITH CC OTITIS MEDIA & URI AGE >17 W/O CC OTITIS MEDIA & URI AGE O-17
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* MEDICARE DATA HAVE BEEN SUPPLEMENTED BY DATA FROM MARYLAND AND MICHIGAN FOR LOW VOLUME DRGS.
** DRGS 469 AND 470 CONTAIN CASES MHICH COULD NOT BE ASSIGNED TO VALID DRGS.
** DRGS 469 AND 470 CONTAIN CASES MHICH COULD NOT BE ASSIGNED TO VALID DRGS.
** TRUE: GEOMETRIC MEAN IS USED ONLY TO DETERMINE PAYMENT FOR DUTLIER AND TRANSFER CASES.
** TRUE THE MEIGHTS ARE BASED ON MEDICARE PATIENT DATA AND MAY NOT BE APPROPRIATE FOR OTHER PATIENTS.

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DUTLIER THRESHOLD 14 15 22 22 9 30	22 22 22 22 28 28 28 28	2222	195 23	22222	82323	23 \$ \$ 2 3 4 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5
GEONETRIC HEAN LOS 3.9 3.1 2.1 12.3	8 4 0 6 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7		6.13	5.2	00000	25.0 25.0 17.4 13.2
ARITHMETIC MEAN LOS 4.3 4.3 5.6	12.6 6.8 10.8 12.8	00000 00000	7.5.5	9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9	5.50 5.50 5.50 5.50 5.50 5.50 5.50 5.50	7.1 5.3 33.3 20.4 15.8
RELATIVE MEIGHTS .6026 .4895 .7404 .3427 3.0258	2.0885 1.0970 1.4817 2.0777 1.3341	1.1032 1.1899 .9698 .5372 1.1451	.7720 1.5691 1.1263 1.2862 .8961	.9448 1.2821 .8264 1.3954 .7571	.9804 .7151 .5744 .7803	.9585 .6625 11.9225 7.3424 5.7811
LARYNGOTRACHEITIS NASAL TRAUMA & DEFORMITY OTHER EAR. NOSE & THROAT DIAGNOSES AGE >17 + OTHER EAR. NOSE & THROAT DIAGNOSES AGE 0-17 HAJOR CHEST PROCEDURES	CONTHER RESP SYSTEM D.R. PROCEDURES W.C. OTHER RESP SYSTEM D.R. PROCEDURES W/O CC PULMONARY EMBOLISM RESPIRATORY INFECTIONS & INFLAMMATIONS AGE >17 WITH CC RESPIRATORY INFECTIONS & INFLAMMATIONS AGE >17 W/O CC	RESPIRATORY INFECTIONS & INFLAMMATIONS AGE 0-17 RESPIRATORY NEOPLASHS MAJOR CHEST TRAUMA WITH CC MAJOR CHEST TRAUMA W/O CC PLEURAL EFFUSION MITH CC	PLEURAL EFFUSION W/O CC PULMONARY EDEMA & RESPIRATORY FAILURE CHRONIC OBSTRUCTIVE PULMONARY DISEASE SIMPLE PNEUMONIA & PLEURISY AGE >17 WITH CC SIMPLE PNEUMONIA & PLEURISY AGE >17 W/O CC	SIMPLE PNEUMONIA & PLEURISY AGE 0-17 INTERSTITIAL LUNG DISEASE WITH CC INTERSTITIAL LUNG DISEASE W/O CC PNEUMOTHORAX WITH CC PNEUMOTHORAX WITH CC	BRONCHITIS & ASTHMA AGE >17 MITH CC BRONCHITIS & ASTHMA AGE >17 M/O CC BRONCHITIS & ASTHMA AGE O-17 RESPIRATORY SIGNS & SYMPTOMS MITH CC RESPIRATORY SIGNS & SYMPTOMS W/O CC	JIHER RESPIRATORY SYSTEM DIAGNOSES MITH CC OTHER RESPIRATORY SYSTEM DIAGNOSES M/O CC G HEART TRANSPLANT G CARDIAC VALVE PROCEDURE M PUMP & M CARDIAC CATH G CARDIAC VALVE PROCEDURE M PUMP & M/O CARDIAC CATH
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~~~~	22277		88 89 90	100000	96 998 1000 1	101

* MEDICARE DATA HAVE BEEN SUPPLEMENTED BY DATA FROM MARYLAND AND MICHIGAN FOR LOW VOLUME DRGS. ** DRGS 469 AND 470 CONTAIN CASES WHICH COULD NOT BE ASSIGNED TO VALID DRGS. NOTE: GEOMETRIC MEAN IS USED DNLY TO DETERMINE PAYMENT FOR OUTLIER AND TRANSFER CASES. NOTE: RELATIVE WEIGHTS ARE BASED ON MEDICARE PATIENTS.

STAY, AND OF WEIGHTING FACTORS, ARITHMETIC AND GEOMETRIC MEAN LENGTH POINTS USED IN THE PROSPECTIVE PAYMENT SYSTEM DIAGNUSIS RELATED GROUPS (DRGS), RELATIVE LENGTH OF STAY OUTLIER CUTDFF UF LIST

				RELATIVE	ARITHMETIC	GEOMETRIC	OUTLIER
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0.001	.00	SUSSE	PASSOCIAN PROCEDURES, W PURP	20,200	13.5	0.44	67
110		2000	ALTHOUGH CANCELL MARKET CONTROLL AND POOL TOTAL	34746	16.3	***	67
011	60	SUKS		3.0/18	1001	16.1	31
1111	0.2	SURG	MAJOR RECONSTRUCTIVE VASCULAR PROC M/O PUMP M/O CC	2.2639	10.6	9,3	27
1115	105	SURG	VASCULAR PROCEDURES EXCEPT MAJOR RECONSTRUCTION M/O PUMP	1,8911	8.2	5.9	24
113	90	SHIRG		2.4590	18.8	14.7	33
114	60	SURG	UPPER LIMB & TOF AMPUTATION FOR CIRC SYSTEM DISORDERS	1.7040	14.0	10.3	28
1115	50	SURG	PERM CARDIAC PACEMAKER IMPLANT W AMI, MEART FAILURE OR SHOCK	4.0516	15.2	12.9	31
116	90	SURG	PERM CARDIAC PACEMAKER IMPLANT W/D AMI, HEART FAILURE OR SHOCK	2.7694	8.3	6.6	25
11.7	90	SURG	SION EXCEPT DEVICE REPLACEMENT	1.2261	6.2	200	-22
118	90	SURG	CARDIAC DEVICE REPLACEMENT	1.7563	4.1	3.0	44
119	90	SURG		.8692	6.3	Mª W	22
150	90	SURG	UTHER CIRCULATORY SYSTEM D.R. PROCEDURES	2.4776	16.6	11.1	58
121	90	MED	CIRCULATORY DISORDERS M AMY & C.V. COMP DISCH ALIVE	1,7162	11.2	9.6	27
155	90	MED	DISORDERS	1.2002	8.7	7.3	25
103	90	MED.	DISORDERS W	1.3979	5.3	2.9	21
124	0.2	MED.	DISORDERS EXCEPT AMI, W CARD	1.1806	6.2	20.0	22
155	62	MED.	CIRCULATORY DISORDERS EXCEPT AMI, W CARD CATH W/O COMPLEX DIAG	*6884	3.2	2.5	10
126	50	MED	ACUTE & SUBACUTE FNORCARDITIS	3.0575	22.7	17.5	15 KT
127		MED.	HEART FAILURE & SHOCK	1.0222	7.9	6.2	24
128	60	MED	DEEP VEIN THROMBOPHLEBITIS	.8513	9.1	8.0	22
129	90	MED.	CARDIAC ARREST, UNEXPLAINED	1.5715	5.9	3.0	21
130	Co	MED	PERIPHERAL VASCULAR DISORDERS WITH CC	.8776	7.8	5.6	24
131	90	MEO	PERIPHERAL VASCULAR DISORDERS W/O CC	. 5862	5.6	0.4	22
132	90	MED	ATHEROSCLEROSTS WITH CC	.7976	0.9	4.7	61
133	97	MED	ATHEROSCLEROSIS W/D CC	.5997	4.6	3.7	14
134	60	MED	41	.6088	5.5	4.4	11
136	5.0	MED	CARDIAC CONGENITAL & VALVULAR DISDRDERS AGE > 17 WITH CC	19221	7.0	5.5	23
136	50	MED	VAL	.6103	60.	3.8	15
137	00	* 034	CARDIAC CONGENITAL & VALVULAR DISCROERS AGE 0+17	.6315	「	100	21
134	5.0	MED	CON 3	.8535	6.2	4	20
1 39	90	MED	ARRHYTHMIA	.5912	4.5	3.6	14
140	50	MED		.6689	2.0	1.1	14
	*					-4	
	THE REAL PROPERTY.						

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LIST OF DIAGNOSIS RELATED GROUPS (DRGS), RELATIVE MEIGHTING FACTORS, ARITHMETIC AND GEOMETRIC MEAN LENGTH OF LENGTH OF STAY OUTLIER CUTOFF POINTS USED IN THE PROSPECTIVE PAYMENT SYSTEM TABLE 5

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00TLIER THRESHOLD 17 13 11 11 11	**************************************	23 23 23 14 16 16 16 17 18	22 22 22 22 22 22 22 22 22 22 22 22 22
GEUMETRIC MEAN LOS 4.4 3.4 3.0 5.8 3.8	112.0 12.0 12.0 12.0 13.0 13.0 13.0 13.0 13.0 13.0 13.0 13	11.0 1.0 1.0 1.0 1.0 1.0 1.0	11.24.64.44.4.4.4.4.4.4.4.4.4.4.4.4.4.4.4.
ARITHMETIC MEAN LOS 5.5 4.2 3.7 7.8 7.8	111111111111111111111111111111111111111	5.45.4 8.8.5.0 0.52.5 1.8.4.5.5 1.8.4.5.5	8 4 5 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8
RELATIVE MEIGHTS -6801 -524 -5500 1-1449	3.4379 2.1344 3.2376 1.8341 2.6797 1.4885 1.5588 1.0566 3.7961	.8382 .9324 .5449 1.1454 .6810 .7541 .7717 .7717 .7717	1.4954 .8651 1.4067 .6689 2.7316 1.1861 .7049 .9878
SYNCOPE & COLLAPSE WITH CC SYNCOPE & COLLAPSE W/O CC CHEST PAIN DITHER CIRCULATORY SYSTEM DIAGNOSES M CC OTHER CIRCULATORY SYSTEM DIAGNOSES M/O CC	RECTAL RESECTION WITH CC RECTAL RESECTION W/O CC RAJOR SMALL & LARGE BOWEL PROCEDURES WITH CC MAJOR SMALL & LARGE BOWEL PROCEDURES WITH CC MAJOR SMALL & LARGE BOWEL PROCEDURES WITH CC PERITONEAL ADMESTOLYSIS W/O CC MINOR SMALL & LARGE BOWEL PROCEDURES WITH CC MINOR SMALL & LARGE BOWEL PROCEDURES W/O CC STOMACH. ESOPHAGEAL & DUODENAL PROCEDURES AGE > 17 WITH CC STOMACH. ESOPHAGEAL & DUODENAL PROCEDURES AGE > 17 WITH CC	* STOMACH. ESOPHAGEAL & DUODENAL PROCEDURES AGE 0-17 ANAL & STOHAL PROCEDURES WITH CC ANAL & STOHAL PROCEDURES WITH CC ANAL & STOHAL PROCEDURES W/O CC HERNIA PROCEDURES EXCEPT INGUINAL & FEMORAL AGE >17 WITH CC HERNIA PROCEDURES EXCEPT INGUINAL & FEMORAL AGE >17 W/O CC INGUINAL & FEMORAL HERNIA PROCEDURES AGE >17 W/O CC INGUINAL & FEMORAL HERNIA PROCEDURES AGE >17 W/O CC HERNIA PROCEDURES AGE 0-17 APPENDECTOMY W COMPLICATED PRINCIPAL DIAG W/O CC	APPENDECTOMY W/D COMPLICATED PRINCIPAL DIAG WITH CC APPENDECTOMY W/D COMPLICATED PRINCIPAL DIAG W/O CC MUUTH PROCEDURES WITH CC MUUTH PROCEDURES WITH CC MUUTH PROCEDURES W/O CC MUTHER DIGESTIVE SYSTEM 0.R. PROCEDURES W/O CC DIGESTIVE MALIGNANCY WITH CC DIGESTIVE MALIGNANCY WITH CC DIGESTIVE MALIGNANCY W/O CC G.I. HEMDRRHAGE W/O CC G.I. HEMDRRHAGE W/O CC
05 KED 05	SURG SURG SURG SURG SURG SURG SURG	SURG SURG SURG SURG SURG SURG SURG SURG	SURG SURG SURG SURG SURG MED MED MED
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MEDICARE DATA HAVE BEEN SUPPLEMENTED BY DATA FROM MARYLAND AND MICHIGAN FOR LOW VOLUME DRGS.

DAGS 469 AND 470 CONTAIN CASES WHICH COULD NOT BE ASSIGNED TO VALID DRGS.

E. GEOMETRIC MEAN IS USED DNLY TO DETERMINE PAYMENT FOR OUTLIER AND TRANSFER CASES.

E. RELATIVE MEIGHTS ARE BASED ON MEDICARE PATIENT DATA AND MAY NOT BE APPROPRIATE FOR OTHER PATIENTS.

DIAGNOSIS RELATED GROUPS (DRGS), RELATIVE WEIGHTING FACTORS, ARITHMETIC AND GEOMETRIC MEAN LENGTH OF STAY, AND
LENGTH OF STAY DUTLIER CUTOFF POINTS USED IN THE PROSPECTIVE PAYMENT SYSTEM H

DUTLIER THRESHOLD 24 18 14 25 25	22122	182251	\$ \$ \$ \$ \$ \$ \$	302 30 21 21 21 21 21 21 21 21 21 21 21 21 21	25448	35825	
GEDMETRIC MEAN LOS 6.1 5.5 4.3 7.1	*****	~~~~~ ~~~~~	1100.0	12.3	8.3.000	112.3862	
ARITHMETIC MEAN LDS 7.8 6.5 5.1 9.1 7.6	***********	80000	12.3	11.22	12.8 9.9 9.9 7.7 9.3	13.4	
RELATIVE MEIGHTS .9964 .7834 .5838 1.0416	. 5415 . 5252 . 5253 . 4223	**************************************	4.6881 3.6252 3.0252 1.6509 2.3854	1.6898 1.8768 1.1152 2.2693 2.4731	2.3933 1.2075 1.00269 1.2132	.6806 .9243 .5816 2.4145 2.1776	
176 06 MED COMPLICATED PEPTIC ULCER WITH CC 177 06 MED UNCOMPLICATED PEPTIC ULCER W/O CC 179 06 MED INFLAMMATORY BOWEL DISEASE 180 06 MED G.I. DBSTRUCTION WITH CC	181 06 MED G.I. DBSTRUCTION W/O CC 182 01 SORDERS AGE >17 WITH CC 182 06 MED ESOPHAGITIS. GASTROENT & MISC DIGEST DISORDERS AGE >17 W/O CC 183 06 MED ESOPHAGITIS. GASTROENT & MISC DIGEST DISORDERS AGE >17 W/O CC 184 06 MED ESOPHAGITIS. GASTROENT & MISC DIGEST DISORDERS AGE 0-17 M/O CC 185 06 MED OFWITH & ORAL DIS EXCEPT EXTRACTIONS & RESTORATIONS. AGE >17	186 06 MED * DENTAL E DRAL DIS EXCEPT EXTRACTIONS & RESTORATIONS, AGE 0-17 187 06 MED DENTAL EXTRACTIONS & RESTORATIONS 188 06 MED OTHER DIGESTIVE SYSTEM DIAGNOSES AGE >17 WITH CC 189 06 MED OTHER DIGESTIVE SYSTEM DIAGNOSES AGE >17 W/O CC 190 06 MED OTHER DIGESTIVE SYSTEM DIAGNOSES AGE >17 W/O CC	191 07 SURG MAJOR PANCREAS. LIVER & SHUNT PROCEDURES 192 07 SURG MINOR PANCREAS. LIVER & SHUNT PROCEDURES 193 07 SURG BILIARY TRACT PROC EXCEPT TOT CHOLECYSTECTOMY WITH CC 194 07 SURG BILIARY TRACT PROC EXCEPT TOT CHOLECYSTECTOMY W/O CC 195 07 SURG TOTAL CHOLECYSTECTOMY W C.D.E. WITH CC	196 07 SURG TOTAL CHOLECYSTECTOMY W C.D.E. W/O CC. 197 07 SURG TOTAL CHOLECYSTECTOMY W/O C.D.E. WITH CC. 198 07 SURG TOTAL CHOLECYSTECTOMY W/O C.D.E. W/O CC. 199 07 SURG HEPATOBILIARY DIAGNOSTIC PROCEDURE FOR MALIGNANCY 200 07 SURG HEPATOBILIARY DIAGNOSTIC PROCEDURE FOR MALIGNANCY	201 07 SURG OTHER HEPATOBILIARY OR PANCREAS O.R. PROCEDURES 202 07 MED CIRRHOSIS & ALCOHOLIC HEPATITIS 203 07 MED MALIGNANCY OF HEPATOBILIARY SYSTEM OR PANCREAS 204 07 MED DISORDERS OF PANCREAS EXCEPT MALIGNANCY 205 07 MED DISORDERS OF LIVER EXCEPT MALIGNANCY 205 07 MED DISORDERS OF LIVER EXCEPT MALIG.CIRR.ALC HEPA WITH CC	206 07 MED DISORDERS OF LIVER EXCEPT MALIG.CIRR.ALC HEPA W/O CC 207 07 MED DISORDERS OF THE BILIARY TRACT WITH CC 208 07 MED DISORDERS OF THE BILIARY TRACT W/O CC 209 08 SURS MAJOR JOINT & LIMB REATTACHMENT PROCEDURES 210 08 SURS MIP & FEMUR PROCEDURES EXCEPT MAJOR JOINT AGE >17 WITH CC	

* MEDICARE JATA HAVE BEEN SUPPLEMENTED BY DATA FROM HARYLAND AND MICHIGAN FOR LOW VOLUME DRGS. ** DRGS 469 AND 470 CONTAIN CASES WHICH COULD NOT BE ASSIGNED TO VALID DRGS. NOTE: GENMETRIC MEAN IS USED UNLY TO DETERMINE PAYMENT FOR OUTLIER AND TRANSFER CASES. NOTE: RELATIVE WEIGHTS ARE BASED ON MEDICARE PATIENT DATA AND MAY NOT BE APPROPRIATE FOR OTHER PATIENTS.

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NAME AND OLOGO COCCO COC	LIST OF DIAGNOSIS RELATED GROUPS (DRGS), RELATIVE  BENOTH OF STAY OUTLIER CUTOFF  BENOTH OUTLIER CUTOFF
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MEDICARE DATA HAVE BEEN SUPPLEMENTED BY DATA FROM MARYLAND AND MICHTGAN FOR LOW VOLUME DRGS.

DRGS 469 AND 470 CONTAIN CASES WHICH COULD NOT BE ASSIGNED TO VALID DRGS.

FE. GEOMETRIC MEAN IS USED ONLY TO DETERMINE PAYMENT FOR DUTLIER AND TRANSFER CASES.

FE. RELATIVE WEIGHTS ARE BASED ON MEDICARE PATIENT DATA AND MAY NOT BE APPROPRIATE FOR OTHER PATIENTS.

TABLE 5.

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* HEDICARE DATA HAVE BEEN SUPPLEMENTED BY DATA FROM MARYLAND AND MICHIGAN FOR LOW VOLUME DRGS. ** DRGS 469 AND 470 CONTAIN CASFS WHICH COULD NOT BE ASSIGNED TO VALID DRGS. **
NOTE: GEOMETRIC MEAN IS USED ONLY TO DETERMINE PAYMENT FOR OUTLIER AND TRANSFER CASES. **
NOTE: RELATIVE WEIGHTS ARE BASED ON MEDICARE PATIENT DATA AND MAY NOT BE APPROPRIATE FOR OTHER PATIENTS.

TABLE 5

DIAGNOSIS RELATED GROUPS (DRGS), RELATIVE WEIGHTING FACTORS, ARITHMETIC AND GEOMETRIC MEAN LENGTH OF STAY, AND
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* MEDICARE DATA HAVE BEEN SUPPLEMENTED BY DATA FROM MARYLAND AND MICHIGAN FOR LOW VOLUME DRGS. ** DRGS 469 AND 470 CONTAIN CASES WHICH COULD NOT BE ASSIGNED TO VALID DRGS. NOTE: GEOMETRIC MEAN IS USFD ONLY TO DETERMINE PAYMENT FOR OUTLIER AND TRANSFER CASES. NOTE: RELATIVE WEIGHTS ARE BASED ON MEDICARE PATIENT DATA AND MAY NOT BE APPROPRIATE FOR OTHER PATIENTS.

TABLE 5

FO GROUPS (DRGS), RELATIVE WEIGHTING FACTORS, ARITHMETIC AND GEOMETRIC MEAN LENGTH OF STAY. AND	
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GEOMETRIC NEAN LOS 6.2 1.8 6.0	9.2 6.8		6.5	3.3	2.6	5.3	11.6	9.00	2.8	4.3 2.1 1.7	5.7	2.7	4.8
ARITHMETIC MEAN LOS 9.1 2.4	6 . 6 . 6	0.0	0.0	7	3.5	7.5	12.8	7.7	3.8	2.8	5.9	80.00	5.8
RELATIVE WEIGHTS 1.2840 .3542 1.0441	1.0230	.6829	.6789	. 4553	. 6431 . 4431 . 2788	.9050	1.9237	1.0774	. \$930	1.0294	1.1302	.9360	.6734
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LIST OF DIAGNOSIS RELATED GROUPS (DRGS), RELATIVE WEIGHTING FACTORS, ARITHHETIC AND GEOMETRIC MEAN LENGTH OF STAY, AND LENGTH OF STAY DUTLIER CUTOFF POINTS USED IN THE PROSPECTIVE PAYMENT SYSTEM

TABLE 5

				RELATIVE WEIGHTS	ARITHMETIC MEAN LOS	GEOMETRIC MEAN LOS	OUTLIER THRESHOLD
351	12	MED	* STERILIZATION, MALE OTHER MALE REPRODUCTIVE SYSTEM DIAGNOSES	.4886	3.9	2.9	14
353	13	SURG		2.2997	15.2	12.8	31
354	13	SURG	NON-OVARIAN/ADNEXAL	1.5482	10.2	1.6	23
355	13	SURG	UTERINE. ADNEXA PROC FOR NON-OVARIAN/ADNEXAL MALIG M/O CC	.9929	122	6.7	13
356	13	SURG	FEMALE REPRODUCTIVE SYSTEM RECONSTRUCTIVE PROCEDURES	.7983	6.3	5.7	14
357	13	SURG	UTERINE & ADNEXA PROC FOR OVARIAN OR ADNEXAL MALIG	2,1591	13.2	11.4	67
358	13	SURG	NON-MALIGNANCY	1.2941	0.6	8.1	20
359	13	SURG	PROC FOR	.9025	9.9	6.3	12
360	13	SURG	VAGINA, CERVIX & VULVA PROCEDURES	1569.	2.5	3.6	19
361	13	SURG	LAPAROSCOPY & INCISIONAL TUBAL INTERRUPTION	.6442	3.9	2.7	13
362	13	SURG	0	\$604.	2.0	1.7	5
363	13	SURG	DEC. CONIZATION E RADIO-IMPLANT, FOR MALIGNANCY	1659.	6.4	3.5	91
364	13	SURG	ox	.4262	2.8	2.2	00
365	13	SURG	DINER FEMALE REPRODUCTIVE SYSTEM D.R. PROCEDURES	1.9060	12.7	9.6	28
366	13	MED	MALIGNANCY. FEMALE REPRODUCTIVE SYSTEM WITH CC	1.0916	9.6	6.1	24
367	13	MED	FEMALE REPRODUCTIVE SYSTEM	.5481	4.9	3.2	18
368	13	MED	FEMALE REPRODUCTIVE SYSTEM	.8308	7.2	5.7	22
369	13	MED	DTHER FEMALE	.4920	4.6	3.3	17
370	14	SURG	CESAREAN SECTION W CC	1.0303	7.7	6.5	19
371	14	SURG	CESAREAN SECTION W/O CC	.7164	5.2	5.0	6
372	14	MED		.4927	4.3	3.4	12
373	14	MED		.3212	2.7	2.4	9
374	14	SURG	DELIVERY M STERILIZATION 6/OR D&C	.5641	3.5	3.1	1
375	14	SURG	* VAGINAL DELIVERY W D.R. PROC EXCEPT STERIL 6/OR DEC	.6817		***	15
376	14	MED	POSTPARTUM & POST ABORTION DIAGNOSES W/O D.R. PROCEDURE	.3520	3.3	2.5	11
317	14	SURG	ABORTION DIAGNOSES	.9882	4.7	2.9	17
378	14	MED	A SAMPLE AND A SAM	1877.	4.4	4.2	6
379	14	MED	THREATENFO ABORTION	.2843	2.8	2.1	6
380	14	MED	ABORTION W/N D&C	.3124	2.5	1.9	80
381	4	SURG	ABORTION W DEC. ASPIRATION CURETTAGE OR HYSTEROTOMY	*3694	1.9	1.6	9
382	14	MED		.1309	1.3	1.2	2
383	14	MED	OTHER ANTEPARTUM DIAGNOSES W MEDICAL COMPLICATIONS	.3964	4.6	3.3	11
384	14	MED	DTHER ANTEPARTUM DIAGNOSES W/D MEDICAL COMPLICATIONS	.3512	3.7	2.6	13
385	15		MEDNATES. DIED OR TRANSFERRED TO ANDTHER ACUTE CARE FACILITY	1.2232	6.0	20.1	77

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TABLE 5

AND .	DUTLIER THRESHOLD 36 31 27 27 25 20	7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	2553	23.52	1222	22.58	51552
IGTH OF STAY	GEOMETRIC MEAN LOS 17.9 13.3 8.6 7.4	48.484	1.86.5.1 5.65.5.2	0.0000	12.5	2.9 2.0 7.1 1.5 1.5	40101
RIC MEAN LEN	ARITHMETIC MEAN LDS 10.7	16.9	2.6 15.8 15.8	14.5	16.5	3.9 10.7 17.8 19.9	44848
ARITHMETIC AND GEOMETRIC MEAN LENGTH OF STAY, AND PROSPECTIVE PAYMENT SYSTEM	RELATIVE WEIGHTS 3.6480 1.8267 1.1571 1.4127 .9416	3.5252 1.5206 1.2250 .7259	.3441 1.0145 1.2115 2.6900	2.0871 .9252 1.5222 .8085	2.7146 1.9999 1.0802 4742	.4919 .3954 1.2385 .8128 3.5067	1.5894 .9346 .9743 .9778
LIST OF DIACNOSIS RELATED GROUPS (DRGS), RELATIVE WEIGHTING FACTORS, ARITHMET LENGTH OF STAY DUTLIER CUTOFF POINTS USED IN THE PROSPECT	386 15 * EXTREME IMMATURITY OR RESPIRATORY DISTRESS SYNOROME, NEONATE 387 15 * PREMATURITY W MAJOR PROBLEMS 388 15 * PREMATURITY W/O MAJOR PROBLEMS 389 15 FULL TERM NEONATE W MAJOR PROBLEMS NEONATE W OTHER SIGNIFICANT PROBLEMS	391 15 ** NORMAL NEMBORN 392 16 SURG SPLENECTOMY AGE >17 393 16 SURG ** SPLENECTOMY AGE 0-17 393 16 SURG ** SPLENECTOMY AGE 0-17 394 16 SURG OTHER 0.R. PROCEDURES OF THE BLOOD AND BLOOD FORMING ORGANS 395 16 MED RED BLOOD CELL DISORDERS AGE >17	397 16 MED RED BLDOD CELL DISORDERS AGE 0-17 397 16 MED COAGULATION DISORDERS 398 16 MED RETICULDENDOTHELIAL & IMMUNITY DISORDERS WITH CC 399 16 MED RETICULDENDOTHELIAL & IMMUNITY DISORDERS W/O CC 400 17 SURG LYMPHOMA & LEUKENIA M MAJOR O.R. PROCEDURE	401 17 SURG LYMPHOMA & NON-ACUTE LEUKEMIA W OTHER D.R. PROC W.C. 402 17 SURG LYMPHOMA & NON-ACUTE LEUKEMIA W OTHER D.R. PROC W/O CC 403 17 MED LYMPHOMA & NON-ACUTE LEUKEMIA W.C. 404 17 MED LYMPHOMA & NON-ACUTE LEUKEMIA W/O GC 405 17 * ACUTE LEUKEMIA W/O MAJOR D.R. PROCEDURE AGE 0-17	406 17 SUR\$ MYELDPROLIF DISORD DR POORLY DIFF NEDPL W MAJ O.R.PROC W CC 407 17 SURG MYELDPROLIF DISORD DR POORLY DIFF NEDPL W MAJ O.R.PROC W/J CC 408 17 SURG MYELDPROLIF DISORD DR POORLY DIFF NEDPL W OTHER O.R.PROC 409 17 MED RADIOTHERAPY	411 17 MED HISTORY OF MALIGNANCY W/O ENDOSCOPY 412 17 MED HISTORY OF MALIGNANCY W ENDOSCOPY 413 17 MED DIHER MYELOPROLIF DIS OR POORLY DIFF NEDPL DIAG WITH CC 414 17 MED DIHER MYELOPROLIF DIS OR POORLY DIFF NEDPL DIAG W/O CC 415 18 SURG 0.8. PROCEDURE FOR INFECTIOUS & PARASITIC DISEASES	416 18 MED SEPTECEMIA AGE 217 417 18 MED SEPTECEMIA AGE 0-17 418 18 MED POSTUPERATIVE & PRIST-TRAUMATIC INFECTIONS 419 18 MED FEVER OF UNKNOWN ORIGIN AGE 217 HITH CC 420 18 MED FEVER OF UNKNOWN ORIGIN AGE 217 H/J CC

* YEDICARE DATA HAVE BEEN SUPPLEMENTED BY DATA FROM MARYLAND AND MICHIGAN FOR LOW VOLUME DRGS. ** DRGS 469 AND 470 CONTAIN CASES WHICH COULD NOT BE ASSIGNED TO VALID DRGS. NOTE: GEOMETRIC MEAN IS USED DNLY TO DETERMINE PAYMENT FOR DUTLIER AND TRANSFER CASES. NOTE: RELATIVE WEIGHTS ARE BASED ON MEDICARE PATIENT DATA AND MAY NOT BE APPROPRIATE FOR DIHER PATIENTS.

LIST OF DIAGNOSIS RELATED GROUPS (DRGS), RELATIVE WEIGHTING FACTORS, ARITHMETIC AND GEOMETRIC MEAN LENGTH OF STAY, AND

TABLE 5

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HE HED	GEDMETRIC NEAN LOS N. 4. 4. 4. 4. 4. 4. 3. 2. 2. 2. 4. 3	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	W W W	13.7	40 4 0 6 40 8 8 8 8	3.59	**************************************
HE HED VIRAL ILLNESS AGE >17 HE HED VIRAL ILLNESS AGE >17 HE HED VIRAL ILLNESS & FEVER OF UNKNOWN DRIGIN AGE 0-17 HE HED OTHER INFECTIONS & PRENITED AGNOSES DIAGNOSES HE HED OTHER INFECTIONS & PRENITED AGNOSES OF HENTAL ILLNESS HE ACUTE ADJUST REACT & DISTURBANCES OF FENTAL ILLNESS HE HED DISTORERS EXCEPT DEPERSORY HED DISTORERS FOR PERSONALITY & IMPUSE COMPROL HED DISTORERS OF PERSONALITY OF HER SYMPT TRY MITH CC ALCARDO MENTAL DISTORER DIAGNOSES ACHOLORIA ADJUSE OR DEFENDENCE, DEFIDX OR OTHER SYMPT TRY MITH CC ALCARDO DESIDE MENTER FOR INJURIES ALCARDO DEPENDENCE, COMBINED REHAB & DEFUX THERAPY ALCARDO DESIDE MENTER OF INJURIES ALCARDON DESIDE MENTER OF INJURINT AND CC	ARITHMETIC MEAN LGS 5.5 5.0 11.2 21.6 6.0	8.3 10.6 13.0 8.3	7 8 8 ° ° ° ° ° ° ° ° ° ° ° ° ° ° ° ° °	13.6 17.6 12.9 15.6	9.00	3.6	*****
18 MED VIRAL ILLNESS AGE >17 19 MED OTHER INFECTIOUS & PARASITIC DISEASES DIAGNOSES 19 MED OTHER INFECTIOUS & PARASITIC DISEASES DIAGNOSES 19 NEG COTHER OF PROPERTY OF PARASITIC DISEASES OF MENTAL ILLNESS 19 NEG COURT OF PARASITIC DISTANCES OF PSYCHOSOCIAL DYSFUL 19 MED DISTANCES EXCEPT OFPRESSIVE 19 MED DISTANCES EXCEPT OFPRESSIVE 19 MED DISTANCES EXCEPT OFPRESSIVE 19 MED DISTANCE OF PRESONALITY & IMPULSE CONTROL 19 MED DISTANCES OF PRESONALITY & IMPULSE CONTROL 19 MED DISTANCE OF PRESONALITY & IMPULSE SUPPRINCE 20 ALCOHOL/DRUG ABUSE OR DEPROBENCE, DEFT AMA 21/2/DRUG DEPROBENCE, COMBINED REHAB & DETOX THERAPY 22 ALC/DRUG DEPROBENCE, COMBINED REHAB & DETOX THERAPY 23 ALC/DRUG DEPROBENCE, COMBINED REHAB & DETOX THERAPY 24 ALC/DRUG DEPROBENCE, COMBINED REHAB & DETOX THERAPY 25 ALC/DRUG DEPROBENCE, COMBINED REHAB & DETOX THERAPY 26 ALC/DRUG DEPROBENCE, COMBINED REHAB & DETOX THERAPY 27 ALC/DRUG DEPROBENCE, COMBINED REHAB & DETOX THERAPY 28 ALC/DRUG DEPROBENCE, COMBINED REHAB & DETOX THERAPY 29 ALC/DRUG DEPROBENCE, COMBINED REHAB & DETOX THERAPY 20 ALC/DRUG DEPROBENCE, COMBINED REHAB & DETOX THERAPY 21 SURG STAIR GREDIES FOR INJURIES M/O CC 21 MED MULTIPLE TRAUMA AGE >17 M/O CC 21 MED ALLERGIC REACTIONS AGE >17 M/O CC 2	RELATIVE .6255 .6255 1.5333 2.2176	. 6586 . 7305 . 8868 . 9329	.4232 .8149 .5903	.9788 1.3306 .0000 1.7523 2.2498	.7185 1.9218 1.2169 .8207	.4796 .4703 .3470 .7922 .4917	. 5907 . 8976 . 5137 . 9067
	18 MED 18 MED 18 MED 19 SURG 19 MED	19 MED DEPRESSIVE NEUROSES 19 MED NEUROSES EXCEPT DEPRESSIVE 19 MED DISORDERS OF PERSONALITY & 19 MED ORGANIC DISTURBANCES & MENT 19 MED PSYCHOSES 19 MED CHILDHOOD MENTAL DISORDERS	19 MED DIHER MENIAL DISURDER DIAGNUSES 20 ALCOHOL/DRUG ABUSE OR DEPENDENCE, LEFT AMA 20 ALC/ORUG ABUSE OR DEPENDENCE, DETOX OR OTHER SYMPT TRT 20 ALC/ORUG ABUSE OR DEPENDENCE, DETOX OR OTHER SYMPT TRT	20 ALC/DRUG 20 ALC/DRUG 20 NO LDNGER 21 SURG SKIN GRAF 21 SURG WDUND DEB	21 SURG HAND PROCEDURES FOR INJURIES 21 SURG OTHER O.R. PROCEDURES FOR INJURIES 21 SURG OTHER O.R. PROCEDURES FOR INJURIES 21 MED MULTIPLE TRAUMA AGE >17 MITH CC 21 MED MULTIPLE TRAUMA AGE >17 M/O CC	21 MED * MULTIPLE TRAUMA AGE 0-17 21 MED ALLERGIC REACTIONS AGE 217 21 MED * ALLERGIC REACTIONS AGE 0-17 21 MED POISONING & TOXIC FFFECTS OF DRUGS AGE 217 21 MED POISONING & TOXIC FFFECTS OF DRUGS AGE 217	21 MED POISONING & TOXIC FFFECTS OF DRUGS A 21 MED COMPLICATIONS OF TREATMENT WITH CC 21 MED COMPLICATIONS OF TREATMENT W/O CC 21 MED JTHER INJURY, POISONING & TOXIC EFF 21 MED OTHER INJURY, POISONING & TOXIC EFF
	22222	3444 E	***	****	*****	4444	4444

** DRGS 469 AND 470 CONTAIN CASES WHICH COULD NOT BE ASSIGNED TO VALID DRGS. ** DRGS 469 AND 470 CONTAIN CASES WHICH COULD NOT BE ASSIGNED TO VALID DRGS. ** DRGS 469 AND 470 CONTAIN CASES WHICH COULD NOT BE ASSIGNED TO VALID DRGS. ** GEOMETRIC MEAN IS USED ONLY TO DETERMINE PAYMENT FOR OUTLIER AND TRANSFER CASES. ** RELATIVE WEIGHTS ARE BASED ON MEDICARE PATIENT DATA AND MAY NOT BE APPROPRIATE FOR OTHER PATIENTS.

MEIGHTING FACTORS, ARITHMETIC AND GEOMETRIC MEAN LENGTH OF STAY, AND POINTS USED IN THE PROSPECTIVE PAYMENT SYSTEM LIST OF DIAGNOSIS RELATED GROUPS (ORGS), RELATIVE LENGTH OF STAY DUILIER CUTOFF

THRESHOLD	23	22	34	28	24	17	32	62	15	0	20	21	28	0	0	30	35	26	51	27	
GEUMETRIC MEAN LOS	4.6	4.1	16.1	10.0	4.9	2.4	13.7	5.3	3.4	1.8	2.8	2.9	6.6	0.	0.	18.1	17.1	8.3	33.3	8.8	
MEAN LOS	10.1	8.5	22.8	14.6	9.5	***	18.3	7.0	4.5	2.2	5.0	8.9	15.5			20.7	29.6	14.8	45.9	12.8	
MEIGHTS	1.9811	2.5317	3,7113	1.7964	1.0495	.7198	1.7517	.7633	.4740	.3172	.5383	.4723	2.4679	00000	0000.	4.0896	10.7296	2.7107	11.8772	3.1757	
	CARE FACILITY			MENT OR OTHER O.R. PROC	URE	DIHER CONTACT W HEALTH SERVICES				S SECONDARY DIAGNOSIS	MALIGNANCY AS SECONDARY DIAGNOSIS	ATUS		ID AS DISCHARGE DIAGNOSIS		ROCS OF LOWER EXTREMITY		DURE AGE >17	ISIS WITH TRACHEDSTONY	ENTILATOR SUPPORT	
	BURNS, TRANSFERRED TO ANOTHER ACUTE CARE FACILITY	EXTENSIVE BURNS M/O D.R. PROCEDURE	NON-EXTENSIVE BURNS W SKIN GRAFT	NON-EXTENSIVE BURNS & MOUND DEBRIDEMENT OR OTHER O.R. PROC	NON-EXTENSIVE BURNS W/O 0.R. PROCEDURE	D.R. PROC M DIAGNOSES OF DIHER CONT		SIGNS & SYMPTOMS W CC	SIGNS & SYMPTOMS W/O CC	AFTERCARE W HISTORY OF MALIGNANCY AS SECONDARY DIAGNOSIS	AFTERCARE W/O HISTORY OF MALIGNANCY	STHER FACTORS INFLUENCING HEALTH STATUS	UNRELATED OPERATING ROOM	PRINCIPAL DIAGNOSIS INVAL	UNGROUPABLE	BILATERAL OR MULTIPLE MAJOR JOINT PROCS OF LOWER EXTREMITY		ACUTE LEUKEMIA M/M MAJOR O.R. PROCE	RESPIRATORY SYSTEM DIAGNOSIS WITH I	RESPIRATORY SYSTEM DIAGNOSIS WITH V	
		MED	SURG	SURG	MED	SURG	MED	MED	MED	MED	MED	MED		**	:	SURG	SURG			MED	
	22	22	22	22	22	23	23	23	23	23	23	23				08	22	17	0.	40	
	456	457	458	459	440	195	462	463	494	465	464	194	468	649	410	471	412	473	474	475	

DRGS 469 AND 470 CONTAIN CASES WHICH COULD NOT BE ASSIGNED TO VALID DRGS.

• GEOMETRIC MEAN IS USED ONLY TO DETERMINE PAYMENT FOR DUTLIER AND TRANSFER CASES.

• RELATIVE WEIGHTS ARE BASED ON MEDICARE PATIENT DATA AND MAY NOT BE APPROPRIATE FOR OTHER PATIENTS. MFDICARE DATA HAVE BEEN SUPPLEMENTED BY DATA FROM MARYLAND AND MICHIGAN FOR LOW VOLUME DRGS.



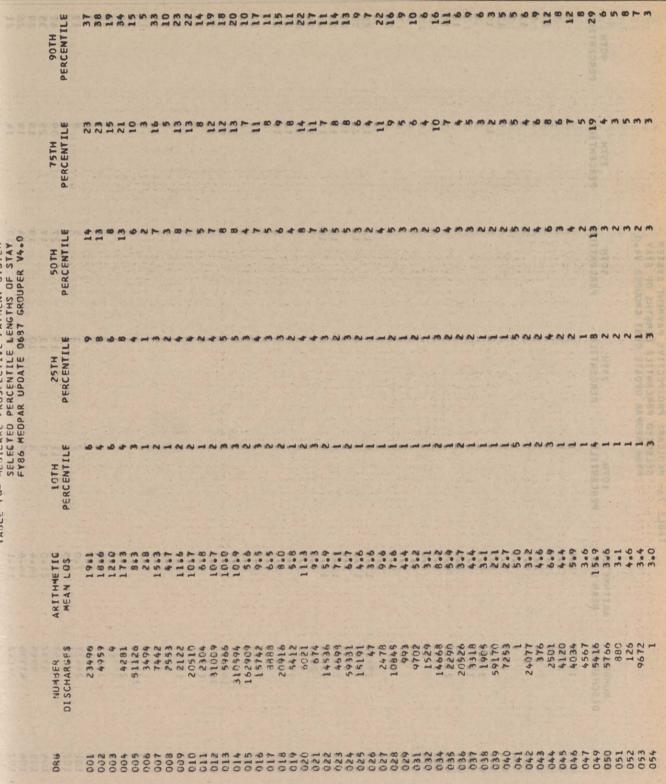


TABLE 7a- MEDICARE PROSPECTIVE PAYMENT SYSTEM

SYSTEM	STAY	0.44
r SY	OF	JPER
MEN	SHL	GROUPER
E PA	LENG	0687
PROSPECTIVE PAYMENT	PERCENTILE LENGTHS	UPDATE
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MEDICARE	SELECTED	FY86
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		PERCENTILE	20	26	0 i	2 2	3.50	72	26	51	m «	31	34	61	14	(C)	13	***	25.	52	151	14	1.1	11	20 .	17		7.2	11	-	0 0		1	151	10	56	07	91	24	15		11	30	18	12	7
		PERCENTILE	01	17	87.	71	22	18	19	01	20 12°	. 200	20	14	11		20 -	3.1	10	11	. 80	6	00		n ?	- =	20	12	4	5	9	D 15		10	-0	50	101	12	16	11	11	6	61	71	9 00	15
ENT SYSTEM	GROUPER V4.0	PERCENTILE	12		12	7 7	11,	11	13	-	•		11	10	80	m ı	n "	700	9	0 00	2	9	4	י חצ		<b>3</b> U	3	27		3	•	* "	- 1	9	+	14	11	10	12	8	80	7	17	- 6	2	S 200 S 200 S
PROSPECTIVE PAYMENT	POATE 0687	PERCENTILE	a	9 6	01		10	9	6	*		v .en	. 0	1	9	- 1	~ "	7.	7	- 0		•	2	m 7	7.		2	72		7	ch c	4.4	2.	*	2			, a	200	9		5		0 8	M	2
ABLE 7a- MEDICARE PROS	FY86 MEDP	PERCENTILE		1	9 1		0 0	*	9	m :	2	. ~		*	3		1	- 3		, ,	-	1		2.	1	, ,	,	27	2	1	2 "	2	. 1	2			0	-	9	*	3	3	,	+ a	2	Day 1 Table
TAB		ARITHMETIC MEAN LOS		12.4	15.5	7.01	18.6	14.2	15.2	3 .	6.3	2 9	16.5	11.3	200	7.	4.0	23.6	2000	0.0	6.2	7.4	5.5	0.9	700	2000	8.4	27.0	6.0	4.0	0.0	1.7	3.00	8.0	2.5	17.5	16.5	0.01	14.1	6.6		7.6		1001	0.99	
		NUMBER	1365	6511	60449	1321	29917	8493	7205	53253	4301	4817	20842	122179	132762	14169	57627	22.00	500070	38392	6693	81254	11540	36274	1186	44013	885		214489	20251	360863	2,000	80651	43128	10636	20097	10500	405451 076k	18492	2590	4846	1562	44363	3948	38318	11317
		DRG	108	109	110	111	113	114	1115	116	111	611	120	121	122	123	124	125	127	128	129	1.30	131	132	133	134	136	137	138	139	140	141	143	144	145	146	147	140	150	151	152	153	45.	155	157	158

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10	STAY	V4.0
SYSTEM		
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2	SELECTED PERCENTILE LENGTHS	FY86 MEDPAR
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MEDICARE PROSPECTIVE PAYMENT		
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75TH PERCENTILE	8	•	,	•	13	0.0		1	•	61	12	. 60	•	•			11	8	•	*	•	80	•		2	9:	42	02	15	2 =	12	8	13	11	13	12	01	11			15
PERCENTILE	\$	•	•		01	80 V	0		2	12		•	5	•			-	5	*		2	2	* ^	· Committee of the comm	3	*!	15	*1	=:	6	6	9	23	0	00	-	91	-	• •	E LOW SOUTH	12
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ARITHMETIC MEAN LOS	6.5	4.5	4.0	4.4	11.5	38.3	8.0	4.9	3.5	15.4	200	6.3	6.9	6.4	5.7	6.4		7.0	6.4	2.0	3.5	9.9	0.4	4.9	4.2	50	19.7	16.4	1104	9.6	10.4	6.9	15.3	12.8	10.0	9.3	8.0	0.6	5.9	4.1	13.8
NUMBER	78212	7204	70711	. 13	2560	1152	1540	7646	2652	12808	1188	2160	152700	10761	11595	6044	7160	69208	8018	108167	54	5455	7946	43761	2845	241	3343	12023	441	1377	76595	+6691	3891	3950	16453	30357	31640	21593	48715	6030	170963
DAG	159	100	161	163	104	165	167	108	169	170	171	173	174	175	176	178	179	180	181	182	184	185	186	188	189	061	192	193	161	196	161	861	661	201	202	203	504	205	202	208	509

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VE PAYMENT SYSTEM E LENGTHS OF STAY 0687 GROUPER V4.0	PERCENTILE	201	•==	8 01	13	9	10 IN	0	0.00	m v			•	m r		•	0 60	2	11 8	- 4		• • •		w 4	2	\$ "	2	2	9.0	•	9	· Harden	M M
MEDICARE PROSPECTIVE PAY SELECTED PERCENTILE LENG FY86 MEDPAR UPDATE 0687	25TH PERCENTILE	0 00	n r «		•		+ 12	~ 1	~ ~	2 "	2	2-	. ~	2	•		2	3	- 5	**	0		) m			2	, -	2	m ~	2	Macenin 5	2	2 2
TABLE Ta- MEDICARI SELECTEI FY86 MEI	PERCENTILE	- 5	m a d	140		m m	+-		1						. 2	~ ~	2	2	**		3	~ ~	-	2 -	2			2	2	-	THE PERSON SHALLING	1 MANUE I	
1	ARITHMETIC MEAN LOS	15.0	14.7	9.7		9.9	5.3	9.4	0.4	1.4	4.4	4.5	7.0	5.9	10.3	5.7	10.5	9.9	10.0	9.0	11.7	1.1	5.2	6.2	5.9	7.0	3.0	2.0	7.1	6.1	6.9	5.8	3.4
	NUMBER	122745	4813	14682	6116	18582	7224	3930	10436	20075	4008	5393	4482	11284	9244	2651	40140	2370	59051	16509	2315	153647	2738	3021	7645	5455	6961	1	32242	10207	37430	7217	1665
	ORG	210	212 213	215	217	218	220	222	223	225	227	228	230	231	233	234	236	237	238	240	242	243	245	246	248	249	251	252	253	256	752	259	260

90TH PERCENTILE	
PERCENTILE	200 2 2 4 1 2 4 1 2 1 2 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0
SOTH PERCENTILE	
25TH PERCENTILE	
PERCENTILE	
ARITHMETIC MEAN LDS	
NUMBER	26489 26689 26689 8694 8634 13762 13762 13762 13762 1629 1629 1629 1629 1629 1629 1629 16
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	10.5	FY86 MED	FY86 MEDPAR UPDATE 0687	0687 GROUPER V4.0		
NUMBER	ARITHMETIC	10TH PERCENTIFE	25TH PERCENTILE	SOTH PERCENTILE	75TH PERCENTILE	90TH PERCENTILE
1388	3.6		2		•	-
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20627	13.5	2	• •	1	12	61
1770	2.6	1	-	1		50
10105	4.00		en e	9 7	10	81
569	8-2	-	7	1	10	*1
10137	6.2	. ~	3	5	1	10
48	5.0	2	3		1	6
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TABLE 7a- MEDICARE PROSPECTIVE PAYMENT SYSTEM
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SYSTEM	SELECTED PERCENTILE LENGTHS OF STAY	FY86 MEDPAR UPDATE 0687 GROUPER VS.O.
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		DISCHARGES	1342	9059	21526	62016	32058	8485	63008	4329	7839	4815	18539	132792	69032	57697	89902	1675	38462	9687	50093	37023	27574	94071	7703	2764	1	442841	361137	60330	44559	43537	10652	15865	5958	33868	13504	7578	1109	4913	12007	1	22904	61113
		ORG	108	601	111	112	113	114	1115	1117	118	119	071	122	123	124	125	171	128	129	130	131	132	133	135	136	137	138	140	141	145	143	145	146	141	241	150	151	152	153	155	156	157	629

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	NUMARK	11162	30598	13	3077	1992	4362	10762	32442	3700	46357	10975	12454	7169	30826	216375	149171	6695	2962	32113	142	5357	9333	3430	9498	49815	3891	2327	16455	30564	18345	5477	32831	170243
	DAG	159	161	163	165	166	168	170	172	173	175	175	178	179	181	182	184	185	1 20	188	061	191	193	100	196		199	200	202	203	205	206	201	500

		90TH PERCENTILE	27	14	82	18	54	22	-	21	**	-0	22	6	6	15	13	13	13	35	12	30	19	13	42	22	11	01	11	13	-	16	01	12	6	1	9
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MEDICARE PROSPECTIVE PAYMENT SYSTEM SELECTED PERCENTILE LENGTHS OF STAY	PAR UPDATE 0687	25TH PERCENTILE	10		6		-	0 4		9	3	2	*	2	2	3	2	-	1			-	\$		•			2		2		,	2	2	* Charles and the same of the	2	2
TABLE 75- MEDICARE SELECTED	FY86 MED	PERCENTILE	-0	3	• •	5	3 6	• •	•	2	2	-	2	-	-		-	-	2	2	2	• 10 10 10 10 10 10 10 10 10 10 10 10 10	0.00	2	. 3	2	2	-	2			2	-		3	7	1 3/04
TA		ARITHMETIC MEAN LOS	16.3	7.2	15.6	10.4	20.7	12.0	5.3	7.0			10.6	4.7	4.4	7.3	0.9	12.9	A 40	14.2	6.6	14.7	10.0	7.0	11.8	7.8	5.8	5.3	5.9	6.8	3.6	8.9	5.4	1.9	5.9	3.7	3.4
		NUMBER DI SCHARGES	72808	14	17492	29465	11197	9200	*	3421	10439	8693	3514	8387	5561	4116	8752	156	7003	9949	40201	8905	59420	8894	2321	13517	11754			3860	7554	15072	22568	10311	25480	5222	2066
		DRG	210	212	412	215	217	218	220	221	223	224	525	227	228	230	231	232	234	235	237	238	239	241	242	543	542		848	250	152	253	254	256	258	260	192

TABLE #5- MEDICARE PROSPECTIVE PAYMENT SYSTEM SELECTED PERCENTILE LENGTHS OF STAY FY86 MEDPAR UPDATE 0687 GROUPER V5.0

PERCENTILE	からのようのからのからのいい ちょうしょう しょうしょうりゅうかい しょうしょうしょうしょうしょうしょうしょうしょうしょうしょうしょうしょうしょうし	133
75TH PERCENTILE	8 3 2 2 1 2 4 2 4 2 5 4 1 8 1 8 4 4 5 6 8 4 6 1 8 5 6 8 1 8 1 8 1 1 1 1 1 1 1 1 1 1 1 1 1 1	
SOTH	N. 3 T. W. W. S. S. C. S. C.	. + 10 m
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PERCENTILE	とわらりょ からをする こうとからの よす こうこう こうそう こうこう こうとう しょうかい いっぱい	212
ARITHMETIC MEAN LOS		1004 500 30.7 50.3
NUMBER DI SCHARGES	2	7887 5844 29346 35163 4123
DRG	00000000000000000000000000000000000000	308 309 310 311 312

RE PROSPECTIVE PAYMENT SYSTEM	SELECTED PERCENTILE LENGTHS OF STAY	FY86 MEDPAR UPDATE 0687 GROUPER V5.0
BLE 75- MEDICARE	SELECTED	FY86 MED
7b-		
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# SELECTED PERCENTILE LENGTHS OF STAY FY86 MEDPAR UPDATE 0687 GROUPER V5-0 TABLE 75- MEDICARE PROSPECTIVE PAYMENT SYSTEM

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The V5.0 Grouper software used to assign FY 1986 cases to DRGs is based on the DRG classifications that will be in effect for discharges occurring in Federal FY 1988. For further information on the DRG classifications, see the final notice of DRG classification Changes published elsewhere in this issue of the Federal Register. TECHNICAL NOTE:

Let n represent the number of discharges classified by DRG and let  $x_1$ ,  $x_2$ , . . . .  $x_n$  Let n represent the ordered LOS values for each DRG set of discharges. For the tth percentile, where p=t/100, let np=j+g where j is the integer part and g is the fractional part of np. The  $t_{th}$  percentile, y, is defined as the LOS value at  $x_{np}$ . Each percentile length of stay (LOS) value was computed according to the following formula:

## $Y = (1-g)x_j + gx_{j+1}$

(See SAS User's Guide: Basics, Version 5 Edition, Cary, NC: SAS Institute Inc., 1985, p. 1186.)

### Appendix A—Regulatory Impact Analysis

A. Introduction

Executive Order (E.O.) 12291 requires us to prepare and publish a final regulatory impact analysis for any regulation that meets one of the E.O. criteria for a "major rule"; that is, that would be likely to result in: an annual effect on the economy of \$100 million or more; a major increase in costs or prices for consumers, individual industries. Federal, State, or local government agencies, or geographic regions; or significant adverse effects on competition, employment, investment, productivity, innovation, or on the ability of United States-based enterprises to compete with foreignbased enterprises in domestic or export markets. In addition, we generally prepare a final regulatory flexibility analysis that is consistent with the Regulatory Flexibility Act (RFA) (5 U.S.C. 601 through 612), unless the Secretary certifies that a regulation will not have a significant economic impact on a substantial number of small entities. For purposes of the RFA, we treat all hospitals as small entities. As we noted in the June 10 proposed rule, it is clear that these changes will affect a substantial number of hospitals and the effects on some will be significant. Therefore, the discussion below, in combination with the rest of this final rule, constitutes a combined regulatory impact analysis and regulatory flexibility analysis in accordance with E.O. 12291 and the RFA. It includes our responses to comments received on the initial analysis published June 10, 1987 at 52 FR 22155.

B. Hospitals Included In and Excluded From the Prospective Payment System

With the enactment of section 9304 of Pub. L. 99-509, which added section 1886(d)(9) to the Act, the 58 acute care hospitals located in urban and rural areas of Puerto Rico will be included in with the approximately 5,700 hospitals that are already operating under the prospective payment system, effective with discharges on or after October 1, 1987. Also, effective with cost reporting periods beginning on or after October 1. 1987, alcohol/drug hospitals and units that have been excluded from the prospective payment system under § 412.22(c) or §§412.25 and 412.32, respectively, of the regulations will begin receiving Medicare prospective payment. Twenty two hospitals and 347 units will be affected by this provision. Only 170 hospitals remain excluded from the prospective payment system under sections 1814(b)(3) and 1886(c) of

the Act (Maryland and New Jersey) or as part of demonstration projects (the Rochester region of New York State).

As of March 31, 1987, 769 Medicare hospitals were excluded from the prospective payment system and continued to be paid on the basis of reasonable cost reimbursement, subject to limits on the rate of their cost increases for FY 1988. These hospitals include psychiatric, rehabilitation, longterm care, and children's hospitals. Another 1,419 psychiatric and rehabilitation units in hospitals subject to the prospective payment system are excluded from prospective payment as of the same date. These units, too, are paid on the basis of reasonable cost reimbursement, subject to limits on the rate of their cost increases.

More than 400 hospitals are being paid on various special bases under the prospective payment system, as required by statute. They include hospitals accorded special treatment as described in our regulations at 42 CFR Part 412, Subpart G, such as: sole community hospitals; cancer treatment and research hospitals that meet certain conditions; and rural referral centers.

C. Inclusion of Puerto Rico Hospitals Under the Prospective Payment System

Using the best data available, we have computed the estimated difference between payments to Puerto Rico hospitals under the rules now in effect (§ 413.40) and under the prospective payment methodology prescribed in the Act. We estimate the combined effect on all Puerto Rico hospitals of implementing prospective payments will be an average payment increase of 5.7 percent over projected payments under the present payment provisions.

In computing this impact, we took into account an estimate of payments for indirect medical education costs and payments to disproportionate share hospitals. To simulate projected payments under the present regulations (that is, under the reasonable cost reimbursement system), we used FY 1988 target payment amounts as an approximation of actual payments. Under payment provisions in effect now. hospitals may receive their actual reasonable costs up to the target amount, plus incentive payments if their actual costs are less than their target amount. Using the target amount as a proxy for actual payments may thus result in a slight understatement of the increase Puerto Rico hospitals may receive under this proposal.

D. Inclusion of Alcohol and Drug Abuse Treatment Hospitals Under the Prospective Payment System

The exclusion of alcohol/drug treatment facilities from the prospective payment system is scheduled to end effective with cost reporting periods beginning on or after October 1, 1987. We have not extended the exclusion beyond that date. On the basis of our research and that of the Alcohol, Drug Abuse and Mental Health Administration (ADAMHA), we have redefined four of the five DRGs into which alcohol or drug abuse cases fall (see the final notice on the DRG classification system published elsewhere in this issue of the Federal Register), and we believe that the reconfigured DRGs and the recalibrated weights will result in equitable payments for alcohol/drug related services to Medicare beneficiaries.

As of March 31, 1987, there were 22 alcohol/drug hospitals and 347 alcohol/drug units located within hospitals already subject to the prospective payment system. These hospitals and units will begin receiving prospective payments for discharges occurring during cost reporting periods beginning on or after October 1, 1987. Because our cost data for these hospitals and units were incomplete at the time of the proposed rule, we were unable to quantify the payment impact of our proposal on these facilities.

Commenters on the proposed regulations for FY 1988 cited the lack of an impact analysis of including currently exempt alcohol/drug abuse hospitals and units in the prospective payment system beginning October 1, 1987. The comment was that such a study should be conducted before any action was taken to end the exclusion. We stated in the June 10, 1987 Federal Register that "because our cost data for these hospitals and units are incomplete, we are unable to quantify the payment impact of our proposal on these facilities." In response to comments, we are now including such an impact analysis. The analysis was accomplished using data from the Hospital Cost Report Information System (HCRIS) on alcohol/drug abuse hospitals/units for FY 1985 and FY 1986.

We estimated what payment to these hospitals would be for FY 1988 under reasonable cost reimbursement, subject to the rate-of-increase limits, and then compared these estimates to what we estimate payment would be under the prospective payment system for FY 1988. The analysis shows that, among 216 alcohol/drug abuse hospitals and units

(we excluded 22 hospitals/units with costs in excess of \$7000 per case or less than \$1000 per case) for which information was available in our data base, estimated prospective payment system payments per case in FY 1988 would be approximately 13 percent less than estimated payments per case under the reasonable cost reimbursement methodology. We estimate the total impact of this change to be a reduction in payments of \$8 million, or an average of \$33,600 per hospital or unit.

In many ways, the results of the impact analysis must be interpreted with caution. As we indicated in the June 10, 1987 Federal Register, we continue to believe our data are incomplete, and thus the results of the analysis must be viewed with the

following points in mind.

We believe that it is inappropriate to draw firm conclusions from comparing estimates of payment under reasonable cost with estimates of prospective payments. This is because hospitals under the reasonable cost system do not have the same incentives for efficiency and productivity as hospitals under the prospective payment system. Data indicating reductions in length of stay since the inception of the prospective payment system demonstrate that the system does have an impact on hospital behavior. Once the currently excluded alcohol/drug hospitals and units are subject to the prospective payment system, we expect them to respond similarly to the system's incentives by increasing their efficiency and productivity, while still continuing to provide high quality care to Medicare beneficiaries.

• The case-mix indexes used to estimate prospective payments are those derived form FY 1985 discharges in our alcohol/drug DRG study file (the MEDPAR file for FY 1985 updated through September 1986), grouped in accordance with the DRG definitions that will be in effect for FY 1988 and the final relative weights published in Table 5 of the Addendum to this notice. The distribution of a hospital's or unit's alcohol/drug cases in FY 1988 may be very different from that in FY 1985; however, we can neither project those differences nor reflect them in this

impact analysis.

• Certain crucial data items are missing from the HCRIS files for some hospitals and units. For example, we are missing such key data as the target amounts and the aggregate reimbursement that would be permitted under § 413.40. Where such data are missing, we have no choice but to assume that a hospital or unit is reimbursed its actual inpatient operating

costs, even though the target ceiling amount might be less. As a result, our estimate of hospitals' payments under the present system may be overstated.

· There are, however, circumstances when the lack of a target amount on the file is appropriate. New hospitals are exempt from the § 413.40 limits for up to three years. This exemption historically has been provided because new hospitals tend to experience high startup costs and relatively low utilization as they establish themselves within a community. The same phenomena may occur in newly established units, but they are not similarly exempt from the limits. Hence, their target amounts, which are based on their actual costs during their first year of operation as a unit excluded from the prospective payment system, may reflect the combined effects of high start-up costs and lower-than-average utilization with no constraining effects from the § 413.40 limits to which they are subsequently

We assume that hospitals and units with target amounts reflected in our data base have been in operation longer than hospitals and units without target amounts and have responded to the § 413.40 limits by constraining their costs so as to remain under the target limits. Conversely, we assume that hospitals and units without reported target amounts have been more recently established and, during their first year of operation as excluded units, had incentives to increase their costs so as to ensure a high target amount for future years.

In fact, when one compares the average operating cost per case of those alcohol/drug hospitals and units with established target amounts to the average operating cost per case of those alcohol/drug hospitals and units without established target amounts, the latter group appears to have costs per case about 20 percent higher than the cost per case of the former group. This supports our hypothesis that the high cost of many of the hospitals and units in the analysis is a temporary phenomenon, occurring in the absence of virtually any limitations on their costs. Thus, we would expect differences between actual costs in FY 1988 and payments under the prospective payment system to be smaller than what we have estimated.

 Some of the cost data are not audited to date. In general, use of unaudited data tends to produce higher estimates of costs than audited data.
 However, we applied no audit adjustment. Consequently, our estimates of reasonable cost reimbursement in FY 1988 would likely be lower if completely audited data were used.

Comment: One commenter recommended, if an impact analysis indicated that the alcohol/drug hospitals and units would receive lower payments under the prospective payment system than under cost reimbursement, that these hospitals and units be held harmless for at least their first year under the prospective payment system.

Response: While our impact analysis projects that prospective payments to alcohol/drug hospitals and units will decline relative to projected payments under cost reimbursement, such an effect is not unlike that which would have been estimated for some groups of general hospitals, had they moved immediately from cost reimbursement to fully national rates under the prospective payment system. However, a four-year phase-in to full national rates has afforded those hospitals the opportunity to make adjustments to their operations so as to bring their costs in line with Medicare's prospective payments. The alcohol/drug hospitals and units, on the other hand, were granted an exclusion of up to four years during which we have conducted extensive analyses so as to refine the alcohol/drug DRG classifications. We believe that, like the transition of shortstay hospitals to full national rates, the exclusion of alcohol/drug hospitals and units has afforded them the opportunity to achieve productivity gains, improve practice patterns, and otherwise adjust their operations in light of the prospective payment rates to which they would be subject once the exclusion ended. In addition, we are concerned that a hold harmless provision as suggested by the commenter is tantamount to extending the exclusion for one more year for those alcohol/drug hospitals and units with costs in excess of their prospective payments while increasing incentive payments for those hospitals and units with costs below their prospective payments. Such a provision could result in increased Medicare expenditures compared to simply extending the exclusion. For these reasons, we do not believe that either a hold harmless provision or a separate phase-in of alcohol/drug hospitals and units is necessary or appropriate.

E. Impact on Excluded Hospitals and Units

As noted above, 769 Medicare hospitals and 1,419 units in hospitals included in the prospective payment system currently are paid on a reasonable cost basis subject to the rate-of-increase ceiling requirement of § 413.40. For cost reporting periods beginning in FY 1988, these hospitals would have their individual target amounts increased by the same factor used to update the prospective payment rate effective for FY 1988. This factor is equal to the projected increase in the hospital market basket less two percent, or an increase of 2.7 percent.

As noted in the proposed rule, the effect this change will have on affected hospitals and units will vary depending on each one's existing relationship of costs per discharge to its target amount, and the relative gains in productivity (efficiency) the hospital or unit is able to achieve. For hospitals and units that incur per discharge costs lower than their target amounts, the primary impact will be to affect the level of additional payments made under § 413.40(c). A hospital may receive additional incentive payments for incurring costs that are less than its target amount, but may not receive payments for costs that exceed the target amount. In general, we expect the inceased ceiling on payments to maintain existing incentives for economy and efficiency experienced by excluded hospitals and units.

### F. Sole Community Hospitals and Rural Referral Centers

At present, about 360 hospitals are receiving payments under § 412.92 based on their status as sole community hospitals. Four of these hospitals have received adjustments to their payment rates as a result of decreases in discharges of five percent or greater. Since the changes we are making to § 412.92 are relatively minor, and will not affect our basic policies regarding

sole community hospitals, we expect no impact on these providers.

We are not making major changes to § 412.96 concerning either the qualifying criteria or payments to rural referral centers. Yet because we are updating the discharge and case-mix criteria, some of the 186 hospitals that now meet these criteria may not meet the new ones, while other hospitals that cannot qualify under the present criteria may qualify under the new criteria. Because we lack data on total discharges, we do not know how many hospitals will either fail or meet the updated criteria. Based on previous experience, the number of hospitals affected either way is small.

### G. Analysis of the Quantifiable Impact of Proposed Changes Affecting Rates and Payment Amounts

### 1. Basis and Methodology of Estimates

The data used in developing the quantitative estimates of changes in payments in Table I, below, are taken from FY 1986 billing data and hospital-specific data for FY 1984. As in previous analyses, we compare the estimated effects of changes for FY 1988 to our estimate of the payment amounts in effect for FY 1987.

We have treated all hospitals in our database as if they had the same cost reporting period; that is, a cost reporting period coinciding with the Federal fiscal year. Our model does not take into account any prospective behavioral changes in response to these changes.

The tables and the discussion that follow reflect our best effort to identify and quantify the effects of the changes being effected through this document. It must be emphasized. however, that as a result of gaps in our data, we are unable to quantify some of the effects.

Nevertheless, we have attempted to improve the accuracy and completeness of our data. One such improvement in the quality of our data is the reclassification of 155 teaching hospitals that had been erroniously classified as non-teaching hospitals.

The analysis that follows examines each of the payment changes separately. That is, all variables except those associated with the provision under examination were held constant so as to display the effects of each provision compared to baseline provisions. Thus, in each of columns 1 through 4, we are comparing estimated FY 1987 payments with the payments that would result if only the specified change were made. (This table has fewer columns than the similar table in the proposed rule because we are not making the proposed changes in outlier payments. which were reflected in column 5 of the proposed rule's table.) The final column (5) displays the combined effects of all the previous analyses, as well as reflecting the FY 1988 update factor (which, giving a 2.7 percent increase across the board, generally has a larger effect than all other changes combined), the budget neutrality factor and the payment adjustment for rural referral centers. Also, the combined effects column captures and reflects certain interactive effects that do not present themselves in the analysis of the individual provisions. This last column is the only one in which the effects of simulated FY 1988 payments are reflected.

BILLING CODE 4120-01-M

TABLE I -- ESTIMATED IMPACT OF THE CHANGES IN THE INPATIENT HOSPITAL PROSPECTIVE PAYMENT SYSTEM

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Number of Hospitals 1/	5414		184	335	404	505	170	205	364	102	498		56	97	346	371	322	591	446	256	162	2767	683	1674	338	72		2647	2045	400	202
	All Hospitals	Urban by Region	New England	Mid Atlantic	South Atlantic	Esst North Central	Esst South Central	West North Central	West South Central	Mountain	Pacific	Rural by Region	New England	Mid Atlantic	South Atlantic	East North Central	East South Central	West North Central	West South Central	Mountain	Pacific	Hrhan Hoenitals	0-99 Beds	100-404 Beds	405-684 Beds	685 + Beds		Rural Hospitals	0-99 Beds	0	170 + Beds

Table 1 - Continued	Number of Hospitals1/	Statutory Blend Change2/ (1)	Statutory Change to Case Weighting 2/ (2)	Reclassification and Recalibration2/	Proposed Revised Wage Index2/ (4)	Combined Effect of All Changes 3/ (5)	
Teaching Status	4362	0.8	0.2	0.0	0.1	3.7	t to had
Resident/Bed Ratio Less than 0.25	868	0.2	-0.2	0.1	0.0	2.3	
Resident/Bed Ratio 0.25 or Greater	184	0.8	-0.3	0.0	0.0	2.9	
Disproportionate Share Hospitals (DSH) No Additional Payments	4218	0.0	0.1	-0.2	0.1	2.4	
Urban DSH 100 Beds or More	878	1.6	-0.3	4.0	0.1	4.2	
Urban DSH fewer than 100 Beds Rural DSH	232	3.0	1.6	0.5	0.0	8.5	
Other Special Status Sole Community Hospitals	326	0.0	6.0	-0.3	0.1	6.7	
Rural Referral Centers (RRCs) Both SCH & RRC Rural fewer than 50 beds	202 21 1233	30.6	-0.2 0.1 2.2	e 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0.2	42.6	
Type of Ownership Voluntary Proprietary Government	3257 775 1358	0.1	0.0	0.0	0001	, e, v,	

" Hospital data base excludes Puerto Rico and alcohol/drug hospitals.

Columns 1 through 4 compare payments incorporating each of the specified FY 1988 changes with estimated payments based on parameters published in the Federal Register September 3, 1986 (51 FR 31454) and November 24, 1986 (51 FR 42229). That is, the comparison is not of simulated FY 1988 rates with FY 1987 rates, but of two sets of FY 1987 rates differing only in the incorporation of the change specified in the column title. 13

This column represents simulated FY 1988 payments compared to estimated FY 1987 payments. It includes the update factor of 2.7 percent required by the Act. This column also reflects the various budget neutrality factors required under the Act as well as interactive effects of the various factors which we are not able to isolate. As a result, the values in this column are not equal to the sum of the four previous columns. 3

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## 2. Summary of Statutory and Wage Index Changes

Columns 1 through 3 of Table I indicate the estimated percent change in payments that would result from each of three statutory changes: transition to a 100 percent national Federal rate; calculation of the average standardized amounts on a discharge-weighted basis rather than on a hospital-weighted basis; and the required annual recalibration of DRG weights. Since the statutory update of 2.7 percent has an across-the-board effect on all hospitals, it is not shown in a separate column, but is merely added into the combined effects shown in column 5.

Our analysis of the statutorily mandated changes remains essentially the same as the one we presented in the proposed rule. The only changes in the final analysis are the result of imporivements in the data.

Column 4 of Table I shows the estimated effects of changes to the wage index, which are described in section III of the preamble. Since we have not altered our policy from what we proposed, the only changes reflected in the final analysis are the result of improvements to the wage index data base that we have made since publication of the proposed rule.

### 3. Combined Effects

In the last column of Table I (column 5) we display the combined effects of the previous four columns plus the effect of the statutorily mandated update factor of 2.7 percent. This column is the only one in which simulated FY 1988 payments are compared to estimated FY 1987 payments.

We must point out that there are interactions that result from the combining of the various separate provisions analyzed in the previous columns and which we are unable to isolate. Thus, the values appearing in column 5 do not represent merely the additive effects of the previous columns plus the update factor and the reduction in the rates for rural referral centers. Note that, generally, the largest changes (other than those attributable to the (update factor) are attributable to the statutory change to a 100 percent national Federal rate.

The greatest change between the final and initial analyses can be attributed to the retention of the present policy on outlier payments. As a result of maintaining the present policy, only urban result, only urban hospitals in the East North Central region are projected to have payment reductions, largely as the result of the statutory transition to a 100 percent Federal payment rate.

Overall, our analysis shows changes in payment policy will increase hospital payments by about 3.1 percent. Urban hospitals will receive an average increase of about 2.7 percent, while rural hospitals will receive, on average, a 4.8 percent increase. Rural hospitals in the East South Central region are projected to receive the largest increase of 7.2 percent. The biggest drop in payments is projected for urban hospitals in the East North Central region. On average, they will receive payment reductions of about 1.3 percent. Among groups of hospitals, rural disproportionate share hospitals and rural hospitals with fewer than 50 beds are projected to receive the largest percentage increases, of 8.5 and 7.6 percent, respectively. Large urban hospitals with over 685 beds are expected to receive the smallest percentage increase of two percent.

Table II presents the projected FY 1988 average payments per case for urban and rural hospitals and for the different categories of hospitals shown in Table I, and compares them with the average estimated per case payments for FY 1987. As such, this table presents in terms of the average dollar amounts paid per discharge the combined effects of the proposed changes presented in Table I. That is, the percentage change in average payments from FY 1987 to FY 1988 equals the percentage changes shown in the last column of Table I.

TABLE II-COMPARISON OF PAYMENT PER CASE, FY 1988 COMPARED TO FY 1987

	Number of hospitals	Averge FY 1987 payment per case	Average FY 1988 payment per case
All Hospitals	5,414	\$4,049	\$4,173
Urban by Region:			BOOK FINE
New England	184	4.672	4.769
Mid Atlantic	335	4.847	5.174
South Atlantic	404	4,056	4.170
East North Central	505	4,700	4.63
East South Central	170	3.733	3,996
West North Central	205	4.530	4.627
West South Central	364	4,056	4,146
Mountain	102	4,491	4,627
Pacific	498	5,116	5.272
Rural by Region:	430	3,110	5,27
New England	56	3,321	3,402
Mid Atlantic	97	3.028	3.088
South Atlantic	346	2,579	2.734
East North Central	371	2,773	2.809
East South Central	322	2,222	2,383
West North Central	591	2,485	2,627
West South Central	446	2,323	2,488
Mountain	256	2,764	2,904
Pacific	162	3.256	3.397
Jrban Hospitals	2.767	4.521	4.645
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0 to 99 Beds	683	3,512	3,693
100 to 404 Beds	1,674	4,258	4,381
405 to 684 Beds	338	4,967	5,080
685+ Beds	72	5,832	5,947

TABLE II—COMPARISON OF PAYMENT PER CASE, FY 1988 COMPARED TO FY 1987—Continued

TO SERVICE THE PROPERTY OF THE	Number of hospitals	Averge FY 1987 payment per case	Average FY 1988 payment per case
Rural Hospitals	2,647	2,598	2,723
0 to 99 Beds	2,045	2,304	2,441
0 to 99 Beds	400	2,642	2,750
170 + Beds	202	3,087	3,209
Teaching Status:	I from the late of		And with the state of the
Non-Teaching	4.362	3.393	3,518
Non-Teaching Resident/Bed Ratio Less than 0.25	868	4,629	4,736
Resident/Bed Ratio 0.25 or Greater	184	6.777	6.970
Disproportionate Share Hospitals (DSH):		and the state of t	
No Additional Payments	4,218	3,759	3,850
No Additional Payments Urban DSH 100 Beds or More	878	4,937	5,142
Urban DSH Fewer than 100 Beds	86	3,743	3,951
Rural DSH	232	2,224	2,41
Other Special Status:	a partire planting	MATERIAL PROPERTY.	STATE OF THE PARTY OF THE
	326	2,815	2.896
Sole Community Hospitals (SCHs)	202	3,237	3,385
Both SCH & RRC	21	3,345	3,441
Rural Fewer than 50 beds.		2,188	2.354
Type of Ownership:			
Voluntary	3,257	4,208	4,314
Proprietary	775	3,680	3,817
Government		3,564	3.769

Appendix B—Final Recommendation of Update Factors for Rates of Payment for Inpatient Hospital Services

Section 1886(e)(4) of the Act, as amended by section 9302(a)(2)(B) of Pub. L. 99-509, required that the secretary, taking into consideration the recommendations of ProPAC, recommend an appropriate update factor for FY 1988, which takes into account amounts necessary for the efficient and effective delivery of medically appropriate and necessary care of high quality. Section 1886(e)(4) of the Act also applies to the target rate-ofincrease limits for hospitals and units excluded from the prospective payment system. (We reiterate that this provision of law requiring recommendations applies to FY 1988 only.)

As required by section 1886(e)(5) of the Act, we published the initial recommended FY 1988 update factors that are provided for under section 1886(e)(4) of the Act. We recommended update factors of 0.75 percent for prospective payment hospitals and 1.9 percent for hospitals excluded from the prospective payment system in a notice published in the Federal Register on June 11, 1987 (52 FR 22386). In recommending these increases, we took into account the requirements in section 1886(e)(4) of the Act. Thus, in that notice, we addressed ProPAC's Recommendations 1 through 5. Also, in that notice, we requested public comment on our recommendations.

Under section 1886(e)(5) of the Act, we are also required to provide a final recommendation of appropriate update factors after consideration of public comments. Accordingly, the purpose of this Appendix is to do so.

We note that although we recommended appropriate update factors, requested and received public comments on these recommendations. and are providing final recommendations, Congress actually prescribed the update factors to be used in FY 1988 in section 1886(b)(3)(B)(i)(II) of the Act, as amended by section 9302(a)(1) of Pub. L. 99-509. That is, as explained in the addendum to the final rule, the update factors for FY 1988 for inpatient hospital services for hospitals under the prospective payment system equals the market basket rate of increase forecasted for FY 1988 minus 2.0 percentage points, or 2.7 percent. This same figure is also the rate of increase in the target rate-of-increase limits for hospitals and units excluded from the prospective payment system.

We received 17 items of correspondence during the public comment period concerning our recommendations and our responses to ProPAC recommendations 1 through 5. After consideration of all the arguments presented, we have decided not to change our proposals. Therefore, we recommend update factors of 0.75 percent for prospective payment hospitals and 1.9 percent for hospitals

and units excluded from the prospective payment system.

Comment: We received one comment that agreed with ProPAC's second recommendation of a higher update factor for rural hospitals in order to reduce the differential in the standardized amounts for urban and rural hospitals to a more reasonable level.

Response: We do not agree that there should be different update factors for the urban and rural standardized amounts. As indicated in our response to ProPAC's recommendation (52 FR 22389), we pointed out that Congress has already taken significant steps to increase payments for rural hospitals relative to urban hospitals. We believe that it ProPAC's recommendation to apply separate update factors for urban and rural hospitals were adopted, it would result in overcompensation to rural hospitals because ProPAC's analysis of the first-year prospective payment system cost experience did not take into account all of the statutorilymandated refinements to the prospective payment system that were not already incorporated into the standardized amounts. For example, ProPAC's analysis did not consider the requirement of section 9302(c) of Pub. L. 99-509 that, effective for discharges in FY 1988, the rates be computed on a discharge-weighted basis rather than a hospital-weighted basis. By narrowing the difference between the urban and rural standardized amounts by more

than three percent, this change has a significant impact on rural hospitals.

Therefore, based on prospective payment system cost experience, we do not support different update factors for urban and rural hospitals as recommended by ProPAC. However, we will continue to study the difference between urban and rural payment rates to determine if additional refinements to the prospective payment system would be warranted.

Comment: Although two commenters agreed that the update factor for excluded hospitals should be different from the update factor for prospective payment hospitals, most commenters stated that our recommended update factors of 0.75 percent for prospective payment hospitals and 1.9 percent for excluded hospitals are too low. The commenters generally expressed concern that the Secretary would implement these recommended update percentages. Some commenters stated that the recommendations of both the Secretary and ProPAC concerning the update percentages were arbitrarily determined and are not adequately supported by quantifiable data and analysis.

Response: Both the Department and ProPAC have maintained that the amounts of the update factors have been based partly on judgments concerning the extent to which hospitals can

increase productivity and reduce their costs. The framework we have established for determining the appropriate update factors was outlined in detail in Appendix B of the June 10, 1985 proposed rule (50 FR 24440). This same framework served as the basis for our recommended update factors for FY 1988

The components of the policy target adjustment factor (PTAF) are difficult to completely quantify individually with existing data sources, and the components of the PTAF, to some extent, represent variables reflecting policy-determined targets. However, judgments about what the appropriate targets should be were also based on our experience with the prospective payment system in determining the extent to which hospitals have responded to the incentives of the system. For example, we have observed a significant decline in hospital lengthof-stay for Medicare patients (a 17 percent reduction) since the beginning of the prospective payment system. This reduction translates into reduced hospital costs that we believe should be accounted for in the PTAF.

In addition, we continue to believe that the initial standardized rates were overstated because they were based on unaudited data. This is evidenced by cost data from the first year of the prospective payment system that indicate that Medicare payments exceeded costs for about 80 percent of all hospitals. Studies conducted by ProPAC using data from first-year prospective payment system cost reports also indicate that the standardized rates would be significantly lower if later, audited cost report data were used.

As we stated in the June 11, 1987 notice, we believe that a policy of steady restraint is warranted so that the Medicare program will continue to benefit from the changes in hospital behavior that have resulted from the prospective payment system. We believe it is appropriate to set the update factor at a level below the projected increase in the hospital market basket, and, in doing so, we have taken into account the requirement under section 1886(e)(4) of the Act that the amounts be high enough to ensure the efficient and effective delivery of medically appropriate and necessary care of high quality.

We believe that our recommended update factors of 0.75 percent for prospective payment hospitals and 1.9 percent for excluded hospitals represent increases in Medicare payments that are adequate to maintain access to high quality care for Medicare beneficiaries.

[FR Doc. 87-19988 Filed 8-27-87; 12:15 pm] BILLING CODE 4120-01-M

### DEPARTMENT OF HEALTH AND HUMAN SERVICES

**Health Care Financing Administration** 

[BERC-410-FN]

Medicare Program; Changes to the DRG Classification System

AGENCY: Health Care Financing Administration (HCFA), HHS. ACTION: Final notice.

summary: This final notice specifies certain changes in the Diagnosis-Related Group (DRG) classification system. It also lists diagnosis and procedures for which new or revised identifying codes (in the coding system of the International Classification of Diseases—9th Edition—Clinical Modification (ICD—9—CM) on which DRG assignments are based) have been approved. This final notice also specifies the changes to the classification of alcohol and drug abuse DRGs and lists the revisions to the surgical hierarchies.

DATE: These classification and coding changes are effective for discharges occurring on or after October 1, 1987.

FOR FURTHER INFORMATION CONTACT: Linda Magno, (301) 594–9343.

### SUPPLEMENTARY INFORMATION:

### I. Background

A. Publication of Proposed Changes

On May 19, 1987 we published a notice in the Federal Register (52 FR 18877) proposing certain changes to the DRG classification system. In that proposed notice we discussed the basic DRG classification system and its relation to the Medicare prospective payment system (PPS), the procedures for changes to the coding system on which the DRG system is based, and the role of the Prospective Payment Assessment Commission (ProPAC) in the analysis of DRGs. Also included in the May 19, 1987 notice was our response to certain recommendations included in ProPAC's April 1, 1987 report to the Secretary. Please refer to that notice for a detailed explanation of the above-mentioned issues.

On June 10, 1987 we also published a proposed rule in the Federal Register (52 FR 22080), which included a proposed reclassification of alcohol and drug abuse DRGs and the revision of surgical hierarchies. Please refer to Section II of that proposed rule (52 FR 22081) for more detailed background of these issues. Since the June 10 proposals also affect DRG classification, we are finalizing those changes and responding

to comments on the proposals in this notice.

B. General

Under the PPS for inpatient hospital services, Medicare payment is made at a predetermined, specific rate for each discharge; that payment varies by the diagnosis-related group (DRG) to which a beneficiary's stay is assigned. Cases are classified into DRGs for payment under the PPS based on the principal diagnosis, any additional diagnoses, and any procedures performed during the stay, as well as age, sex, and discharge status of the patient. The diagnostic and procedure information is expressed by the hospital using codes from the International Classification of Diseases, Ninth Edition, Clinical Modification (ICD-9-CM). The intermediary enters the information into its claims system and subjects it to a series of automated screens called the Medicare Code Editor (MCE). These screens are designed to identify cases that require further review before classification into a DRG can be accomplished. After screening through the MCE and any further development of the claims, cases are classified by a computer program called the Grouper into the appropriate DRG.

The DRGs are organized into 23 major diagnostic categories (MDCs), most of which are based on a particular organ system of the body and the remainder of which involve multiple organ systems (such as MDC 18, Infections and Parasitic Diseases, Systemic or Unspecified Sites; and MDC 22, Burns). Accordingly, the principal diagnosis determines MDC assignment. Within most MDCs, cases are then divided into surgical DRGs (based on a surgical hierarchy that orders individual procedures or groups of procedures by resource intensity) and medical DRGs. The medical DRGs generally are differentiated on the basis of diagnosis. Both medical and surgical DRGs may be further differentiated based on, age, discharge status, and presence or absence of complications or comorbidities (hereafter CC). With some exceptions, the Grouper does not consider other procedures, such as nonsurgical procedures or minor surgical procedures that generally do not require use of an operating room (OR). For ease of reference, when multiple DRGs are hereafter referred to in this final notice we will refer to the DRG title and category but we will not specify age and/or CC breaks, respectively. For example; reference to DRGs 277-279 will indicate (Cellulitis) for all three DRGs in lieu of DRGs 277 (Cellulitis age over 69 and/or CC), DRG 278 (Cellulitis age 1869 without CC), and DRG 279 (Cellulitis age 0-17).

### II. Comments on the Proposed Notice

In response to the May 19, 1987 proposed notice, we received 54 timely public comments and in response to the June 10, 1987 proposed notice we received 111 timely public comments. Both notices received comments from representatives from health care associations, hospitals, ProPAC, physicians and physician associations. Additionally, the May 19, 1987 notice received comments from medical record administrators. ProPAC incorporated by reference its recommendations on DRG classifications included in its April 1, 1987 report to the Secretary.

In addition to comments related to each of the proposed DRG classification changes discussed below, we received some comments of a general nature, as follows:

Comment: Two commenters wrote expressing concern with the lack of detail presented in the proposed notice. The commenters recommended that the notice provide full disclosure of the data, methodology, criteria, calculations, supporting documentation and underlying assumptions made by HCFA in reaching its conclusions. One of the commenters recommended that no further DRG changes, including those proposed for FY 1988, be made until criteria for reclassification are developed and published for public comment.

Response: As we have stated previously, we do not believe the rulemaking process requires, nor are most members of the public interested in, the level of detail requested by these commenters. The volume of data used during our evaluation of DRG issues prohibits publication in the Federal Register. Moreover, as evidence of the fact that most readers would find this level of detail burdensome, we note that we received fewer than 40 requests for copies of the diagnosis-specific material concerning the proposed refinements to the CC list. In light of constraints on Federal agency spending, we believe it would be very imprudent to reproduce thousands of pages in the Federal Register. Providing a detailed description of the analytic bases for our proposed changes is a reasonable alternative and is consistent with the level of detail most readers desire. For those individuals wishing more information, a contact person's name and telephone number are published in each proposed and final notice.

We note, in addition, that the Medicare Provider Analysis and Review

(MEDPAR) file of Medicare discharges, which serves as the basic source of data for analysis of DRG classification changes, can be purchased from our Bureau of Data Management and Strategy. (We regret any confusion created by our reference in the May 19 proposed notice to the PATBILL as our source file for our analysis. The MEDPAR file contains the same data as the PATBILL file but is in a simplified, reformatted record layout. Both files contain the same diagnostic and procedure data for up to five diagnoses and three procedures 100 percent of Medicare inpatient hospital bills. Although we use the two names interchangeably, technically we use the MEDPAR file). Other background information and data used in our DRG reclassification efforts are available upon request, including requests under the Freedom of Information Act. At least one of the above-cited commenters has secured information in this manner.

With regard to the comments recommending that detailed criteria for classification be developed and published for comment, we do not believe such steps are necessary. The DRGs, as originally developed, were intended to represent groups of hospital patients who were clinically similar to one another and were relatively homogeneous with respect to resource use as measured by length of stay (LOS). The algorithm used to define the DRGs was designed to establish partitions that would both reduce the variance with respect to length of stay within groups and maximize the differences between groups. It was originally thought that in order to be manageable, the DRGs should number something less than 500.

In our efforts to refine the DRG classifications and respond to changing medical practice, we have attempted to adhere to those principles. Since we have based DRG weighting factors on total charges (standardized to account for variations among hospitals in area wages, teaching intensity, and the proportion of low income patients), we have used standardized charges rather than the length of stay as our primary measure of relative resource use when examining the extent to which a proposed classification change makes the DRGs more or less homogeneous. In addition, with respect to new technologies, we prefer adding cases involving new technologies to an existing DRG, clinical heterogeneity notwithstanding, until there is relatively compelling evidence (based on Medicare patient experience) that a separate DRG would improve both the clinical coherence and the homogeneity

with respect to resource use for the new DRG. Moreover, we described the bases on which we proposed DRG classification changes in our proposed notice. In general, the changes we have proposed are ones that either reduced within DRG variance among patients in resource use (as measured by standardized charges); incorporated new ICD-9-CM codes into the DRGs; incorporated new technologies into the DRGs; or represented housekeeping changes, such as consistency or other logic checks. In the case of the CC refinement included in the May 19 notice, the proposed exclusions were based on clinical review using the principles stated in that notice.

Generalizing beyond these goals to overriding criteria could be counterproductive in that the criteria thus adopted may be to narrow to permit adoption of a reasonable DRG classification change or too broad to forestall consideration of reassignment of each and every ICD-9-CM code from the DRG(s) to which it is presently assigned to all other possible DRGs. We believe it is better to continue to evaluate each DRG classification issue independently and that it is sufficient to describe the analytic basis upon which we propose each of the individual DRG

classification changes.

We will continue to base our decisions on clinical grounds, comparability of the average charge for one type of case to the mean for the DRG in which it is classified and the DRG to which its movement is proposed, frequency of the procedure or diagnosis at issue, variation in a particular DRG relative to DRGs in general, and other issues pertinent to the type of case being considered for reclassification. We believe such individual consideration is superior to the development of criteria that could potentially prevent movement of cases in instances where reclassification is appropriate.

Comment: Several commenters raised DRG classification issues that had not been discussed in the proposed notice and requested we make DRG changes. Among the issues raised were the

following:

(1) Analysis of the importance of CC in DRGs not currently partitioned by presence or absence of CC:

(2) Scraping of cornea for smear and culture in infected corneal ulcer cases:

(3) Carcinoma of the mouth; and

(4) Classification of trauma cases. Response: To consider new issues that arise during a comment period. especially those that are not directly related to proposed changes, would require us to make hasty decisions

without the benefit of detailed, reasoned analysis or public comments. Consequently, we do not intend to make it a general practice to make DRG changes, other than those directly related to our proposals, in the final DRG notices.

We will, however, place the issues raised on our agenda for study during FY 1988. We welcome further suggestions for issues to study and encourage commenters to submit detailed proposals early in the Federal fiscal year so that we are not hampered in our decision making process by time constraints imposed by the statutorily required publication process. That is, in order to meet the new statutory requirement for publication of a proposed notice by May 1 of each year, we must complete our evaluative process by no later than March of each year. Suggestions may be submitted to: Grouper Changes, P.O. Box 26681, Baltimore, Maryland 21207.

Comment: One commenter, representing a major health insurer, wrote expressing concern with the increase in the price of Grouper software for FY 1988, and believes that there is no justification for a significant price increase in software developed under a grant from HCFA. Another commenter complained of the price of the DRG Definitions Manual.

Response: The development and distribution of Grouper software is currently handled by Health Systems International (HSI) under a contract (not a grant) with HCFA. Under the contract, HCFA provides specifications on DRG classification changes and decides on whether to accept or reject recommended changes in DRG assignment. The contractor then works with HCFA in order to establish the precise logic that will determine DRG assignment, modifies the Grouper software accordingly, and performs extensive quality controls to ensure accuracy of DRG assignment. The Medicare fiscal intermediaries (FIs) must purchase the Grouper software from HSI to process and pay inpatient hospital claims under the Medicare prospective payment system. HSI is also obligated to provide documentation on the Grouper and Medicare Code Editor and to furnish a DRG Definitions Manual to the FIs. Generally, under the terms of HSI's contract, the material is made available to the FIs at cost. Alternatively, HSI itself may defray the cost of equipping the FIs with the software, using funds available under its

HCFA's contract with HSI is silent with respect to HSI's operations in the

private sector. HSI is neither required to market Grouper software and the DRG Definitions Manual nor is it prohibited from doing so. Therefore, HSI is free to market its products in a manner similar to any other business. We take this approach in order to promote efficiency and competition in the marketplace. In this connection, we note that HSI is not the sole source of Grouper software; the software is also programmed and marketed by several other firms, and HCFA makes these materials available to the public through the National Technical Information Service. If a prospective purchaser of Grouper software is dissatisfied with the price charged by one source, it is free to bargain with that source or to seek the material elsewhere.

HSI advised us that the original price of the Grouper software was set in 1982 based on cost estimates furnished by Yale University, the developer of the original DRG system, and that 1987 marks the first increase in the price of the software since 1982.

Comment: One commenter suggested that we provide information on the distribution of cases before and after reclassification changes.

Response: In light of the nature and number of DRG classification changes we are adopting for FY 1988, we concur with the commenter and refer the reader to Tables 7a and 7b of our final notice of changes to the inpatient prospective payments system and FY 1988 rates. The FY 1986 Medicare discharges used for recalibration of the DRG relative weights were grouped in accordance with both the current (FY 1987) DRG classifications for Table 7a and the revised DRG classifications we are adopting with this notice for Table 7b. Both tables show the number of cases and the arithmetic mean length of stay for cases in each DRG as well as the lengths of stay for cases at the 10th. 25th, 50th, 75th and 90th percentiles within the distribution for each DRG.

Comment: One organization recommended that the revised Grouper software be made available simultaneous with publication of the proposed notice of DRG classification changes to permit the public the opportunity to test the effects of the proposed changes on data more current than that available to and used by HCFA for reclassification and recalibration.

Response: In order to incorporate as many changes as possible into each year's proposed notice of DRG classification changes, we have either extended the period of time over which we conduct analyses (as was done this

year) or, as in 1986, proposed changes in both a DRG classification notice (51 FR 18762) and in the proposed notice of changes to the inpatient hospital prospective payment system and FY 1987 rates ( 51 FR 19970). In either situation, we do not have a revised Grouper program until after the notice(s) of proposed classification changes is published. Rather, we simulate the Grouper revisions in order to analyze the effects of classification changes and, based on those analyses, propose DRG revisions. We acknowledge that this process may involve substantial internal programming and keying of codes to mimic the actual Grouper software, but we find that it generally works adequately to develop revised classifications and weighting factors. The commenters' concern that the final DRG weighting factors would be markedly different from the proposed weighting factors is not borne out by a comparison of weights in Table 5 between the proposed and final notices of changes to the inpatient hospital prospective payment system and FY 1988 rates. While most of the DRG weighting factors vary somewhat, we believe this movement has more to do with the use of a more recent update of the MEDPAR file than with differences in the classification of cases between the proposed and final notices resulting from the use of different software to group cases. The final recalibrated DRG weights are based on nearly 9.7 million Medicare discharges in FY 1986 received in HCFA through June 1987, while the proposed weights were based on 9.4 million records received through February 1987. (The June 10 proposed rule erroneously indicated that approximately 9.5 million bills had been used in recalibration.)

Because the approach we currently use works and based on the specificity of information we publish in the proposed notice, can be replicated by other parties, we believe it is preferable to analyze as many DRG classification changes as possible before publishing our proposed notice. Even if we publish our notice of proposed DRG changes earlier in the calendar year as we did in 1986, the need to consider the recommendations of ProPAC on a timely basis would render obsolete any Grouper based on such earlier notice if we accept recommendations of ProPAC on DRG classification changes other than those addressed in our earlier notice.

III. Proposed Changes and Comments Affecting a Particular MDC ¹

A. MDC 3: Diseases and Disorders of the Eye, Nose, and Throat

We stated in our proposed notice that claims for cochlear implants will continue to be assigned to DRG 49, (Major Head and Neck Procedures).

Comment: One manufacturer of cochlear implants, their congressional representatives, and ProPAC wrote to express disappointment in our decision to continue to assign cochlear implant cases to DRG 49.2 They believe that payment at the DRG 49 rate understates the cost of the procedure and thereby adversely affects access to the device for Medicare beneficiaries. They urged us to adopt ProPAC's recommendation to create a unique DRG for implantation of the device.

Another manufacturer of the device wrote expressing satisfaction with our interim assignment of cochlear implants to DRG 49, stating that it has been their experience that current Medicare payments based on the DRG 49 weighting factor have been equitable. Noting that cochlear implant technology had not vet stabilized and that future reclassification might be necessary, this commenter observed that the ability to identify and retrieve from Medicare program files data on the use and cost of cochlear implants was of major importance, but a goal already met by the establishment of unique ICD-9-CM codes for the implantation of the device.

Response: As we stated in our proposed notice, Medicare data indicate that the charges for cochlear implants furnished to Medicare beneficiaries are not significantly different from charges for other cases assigned to DRG 49. Consequently, the creation of a new DRG for the procedure at this time would result in a DRG with very few cases and with a weight nearly identical to that of DRG 49. Indeed, based on partial FY 1987 Medicare billing data more recent than that which was available when we prepared our May 19 notice, we find that the average standardized charge for cochlear implant cases furnished to Medicare patients is actually slightly less than for other cases assigned to DRG 49. Thus, adoption of the ProPAC recommendation would actually reduce payment for the implants. Accordingly,

¹ Unless otherwise noted, these changes were proposed in the May 19, 1987 notice.

² For full DRG titles, see Table 5 in the final rule setting forth FY 1988 PPS changes and rates, elsewhere in this issue of the Federal Register.

we find no compelling reason to create such a new DRG at this time.

We are aware that cochlear implant charges may vary from one maufacturer to another and between single channel and multi-channel devices. However, the DRG system was not designed to recognize individual products by manufacturer or model.

We also acknowledge that other data sources, such as survey data gathered by one manufacturer, may indicate that charges for cochlear implant cases are higher than charges for other cases assigned to DRG 49. We note that the survey data submitted to us had not been standardized to adjust for hospital variations in wages, teaching, and proportion of low-income patients. Moreover, we believe that, wherever possible, it is appropriate to base the Medicare DRG weighting factor on data exclusively from Medicare patients to ensure consistency and comparability of types and sources of data used for all DRGs.

Finally, as we have previously stated, we view the classification of cochlear implant cases to DRG 49 as an interim measure while we continue to study the issue on the basis of more complete data reflective of Medicare beneficiary utilization of cochlear implants and their estimated cost. This will be facilitated by the establishment of a new unique set of codes for cochlear implants. (These codes became effective October 1, 1986.) Should reclassification of cochlear implants then prove appropriate, we will consider it at that time.

B. MDC 4: Diseases and Disorders of the Respiratory System

Based on numerous comments and our analysis of FY 1985 MEDPAR data, we determined that significantly more hospital resources were used to treat patients requiring mechanical ventilation than other patients with a principal diagnosis within the respiratory system MDC. We also observed that while ventilator patients were dispersed throughout the respiratory system DRGs, that they were more comparable to each other than to other cases in MDC 4. Finally, among ventilator patients, substantial differences in resource use also were found to be related to whether ventilator access was achieved through endotrachial intubation or tracheostomy.

Accordingly, we proposed to create two new interim DRGs for MDC 4. Cases presenting a principal diagnosis in MDC 4 and one of the tracheostomy procedure codes [31.1, Temporary tracheostomy; 31.21, Mediastinal tracheostomy; 31.29 Other permanent tracheostomy) would be assigned to a new DRG 474 (Respiratory System Diagnosis with Tracheostomy). This DRG would be ordered above all other DRGs in MDC 4, including the surgical DRGs.

We also proposed creation of a new DRG for cases involving mechanical ventilation through endotrachial intubation. A new medical DRG 475 (Respiratory System Diagnosis with Ventilator Support), would be established for cases presenting a principal diagnosis assigned to MDC 4 and showing both non-OR procedure codes 93.92, Other mechanical assistance to respiration, and 96.04, Insertion of endotrachial tube.

Comment: The American Association for Respiratory Care, the American College of Chest Physicians, the National Association of Medical Directors of Respiratory Care, ProPAC, and numerous other commenters wrote to express general support for the creation of DRGs 474 and 475. In addition, several of the commenters encouraged expansion of this proposal to patients with other than respiratory diagnoses.

Response: We appreciate the support of these commenters as we continue to refine the DRG classification system. As we stated in our proposed notice, we will continue our research in this area, including analysis of superior means of identifying ventilator cases and ways to address this issue in post-surgical cases and/or patients with non-respiratory principal diagnoses.

Comment: One commenter found our expressed concern with the potential for abuse of the proposed new DRGs 474 and 475 offensive, but the National Association of Medical Directors of Respiratory Care and ProPAC shared our concern that the new DRGs may create financial incentives for hospitals to pressure physicians to intubate patients or perform tracheostomies.

Response: We regret that anyone took offense at the cautionary statement included in the preamble of the proposed notice concerning the possibility that changes in medical practice or the reporting of such practices may be precipitated by the new DRGs. We did not intend to cast aspersions upon the medical community or the ethics of physicians or hospital personnel. However, we recognize that there may be an occasional questionable situation in which DRG classification may influence the course of patient treatment or the reporting of the treatment provided. Indeed, as the basis for their recommendations for DRG classification changes, many

commenters routinely claim that the DRG definitions and weighting factors affect medical practice patterns, limit Medicare beneficiary access to the most up-to-date and sophisticated medical technologies, and subject physicians to financially-motivated pressure by hospital managers. If we are to believe that failure of the DRGs to provide higher payment for cases involving certain technologies discourages their use, we may reasonably anticipate that the recognition of procedures and technologies such as tracheostomies and mechanical ventilation in relatively high-weighted DRGs may encourage their use. Further, scattered instances of program abuse do occur and we are required by law to attempt to discover such abuses and institute corrective action.

Finally, we believe it is in the public interest to advise the medical community of our intent to target DRGs 474 and 475 for medical review by the PROs to ensure that use of the diagnoses and procedures that result in assignment of cases to these DRGs is medically reasonable and appropriate.

Comment: One commenter noted that the proposed notice stated that DRG 474 would be ordered above all other DRGs in MDC 4 but was silent as to the ordering of DRG 475. She requested clarification of this point.

Response: Because it contains both OR and non-OR procedures, DRG 474, is not characterized as either a medical or a surgical DRG. Nevertheless, it is ordered first in the hierarchy for MDC 4 and, thus, precedes all surgical and medical DRGs in the MDC. DRG 475, will be assigned to cases with a respiratory system principal diagnosis when neither a temporary tracheostomy nor any operating room procedure is performed and both Endotracheal intubation (code 96.04) and Other mechanical assistance to respiration (code 93.92) are performed. Medical DRGs generally are differentiated by principal diagnosis, so that there is usually no hierarchy beyond the surgical DRGs because the groups of diagnoses are mutually exclusive. However, while DRG 475 is technically a medical DRG. it is not specific to a subset of diagnoses within MDC 4 but rather can be assigned to a case with any principal diagnosis in MDC 4 if the specified procedures are performed. Because DRG 475 is procedure-driven, it is ordered below the surgical DRGs and above the medical DRGs in the MDC.

Comment: One commenter expressed concern that the DRG system does not adequately take into consideration patients who develop acute respiratory

failure. The commenter believes there are well-defined, medically acceptable definitions of acute respiratory failure that should be integrated into the DRG

Response: We believe the changes in the coding of respiratory failure and the creation of new DRGs 474 and 475 for patients requiring ventilator support will adequately address the issue of patients developing respiratory failure. A new ICD-9-CM diagnosis code, 518.81, Respiratory failure, has been adopted and will become effective October 1, 1987. Unlike the previous code for respiratory failure (799.1), this new code is not in Chapter 16 (Symptoms, Signs, and Ill-Defined Conditions) of the ICD-9-CM; therefore, hospitals will now be permitted to code respiratory failure as a principal diagnosis when it is the reason for admission.

### C. MDC 5: Diseases and Disorders of the Circulatory System

### 1. Major Vessel Resection With Replacement

We proposed to assign procedure codes 38.48, Abdominal artery resection with replacement, 38.47, Abdominal vein resection with replacement, and 38.48 Lower limb artery resection with replacement, to DRGs 110 and 111 Major Reconstruction Vascular Procedure without Pump), and remove them from DRG 112 (Vascular Procedures except major reconstruction. without pump). We did not receive any comments on this proposal, so it will be adopted without change.

### 2. Malignant Hypertension

We proposed to continue to classify malignant and benign types of hypertension disease into DRG 134

(Hypertension).

Comment: One commenter disagreed with our decision concerning malignant hypertension. The commenter noted that, in his experience, hypertension is rarely coded as the principal diagnosis without another acute diagnosis. The commenter believes that further study is needed on the conditions and charges in this category before a decision is made.

Response: While we agree in principle with the commenter that further study may be warranted, we note that the comment is generally non-specific as to other diagnoses that, in conjunction with malignant hypertension, account for variation in resource use. In the FY 1985 MEDPAR file on which our analysis was based, the vast majority (93 percent) of malignant hypertension cases were coded 401.0 and had standardized charges that were greater than the mean charges for the DRG by only \$100 (less

than 4 percent). The 208 cases involving hypertensive heart and renal disease (code 404.0), while considerably more costly than the average case in DRG 134, comprised less than 1.6 percent of all malignant hypertension cases and less than one-half of one percent of all cases in the DRG. In light of these findings, we are not prepared to reclassify malignant hypertension cases at this time.

In addition, we suggest that commenters concerned with the DRG classification of such cases furnish to us additional information on classification problems in DRG 134 by writing to the Grouper Changes address published in section II. of this notice. We are not persuaded that further analysis, uninformed by more specific information, would be fruitful at this

### 3. Acute Myocardial Infarction

We proposed to continue to classify acute myocardial infarctions (AMI) into DRGs 121. (Circulatory Disorders with AMI and Cardiovascular Compensation Discharged Alive) 122, (Circulatory Disorders with AMI Without Cardiovascular Compensation Discharged Alive) and 123 (Circulatory Disorders with AMI, Expired).

Comment: Two commenters urged that we reconsider our decision not to reclassify AMI cases at this time. The commenters believe the number of cases subject to misclassification due to coding guidelines is small and should not cause us to postpone quick action on the issue of reclassification of AMI cases. One commenter also suggested that the ICD-9-CM Coordination and Maintenance Committee "develop more ethical AMI coding guidelines.'

Response: We believe that it is premature to revise the classification of AMI cases at this time. As we pointed out in our proposed notice, the current ICD-9-CM coding guidelines can result in the assignment to DRGs 121-123 of cases that do not involve a current, documented AMI but rather involve the admission of patients who had a myocardial infarction within the preceding eight weeks. Such patients may be hospitalized for recurrence of symptoms without an AMI, for regulation of their medication, or for further evaluation of their condition, including cardiac catheterization to determine the need for coronary bypass surgery. As many as 25 percent of FY 1984 AMI cases among Medicare beneficiaries discharged alive show a length of stay of seven days or less. Physicians advise us that the generally recognized standard of care for patients admitted with a documented AMI is at least seven days. Accordingly, cases

with lengths of stay of less than seven days may in fact represent cases not admitted for the treatment of an AMI but nonetheless classified as AMIs because of the eight-week rule. We believe the commingling of such cases with cases admitted and treated for documented AMIs is responsible in part for the variation in resources in these

We appreciate the commenters support for changes to the coding guidelines. The ICD-9-CM Coordination and Maintenance Committee addressed coding of AMIs at its July 1987 meeting. The National Center for Health Statistics solicited comments and announced plans to propose a revision at the November 1987 meeting of the ICD-9-CM Coordination and Maintenance Committee. We will continue to work with the ICD-9-CM Coordination and Maintenance Committee on the coding issue. We encourage commenters to address suggestions for revision of the AMI diagnosis code to:

Ms. Sue Meads, Co-Chairperson, ICD-9-CM Coordination-Maintenance Committee, National Center for Health Statistics, Room 2-19 Center Building, 3700 East-West Highway, Hyattsville, Maryland 20782.

In light of the possibility of coding changes that may affect the assignment of cases currently in the AMI DRGs, we believe it is more appropriate to delay action at this time. In general, we believe that making changes in a DRG that we expect to have to revise further a year or two later based on coding changes would create confusion that is likely to outweigh the potential benefits of the classification change.

In addition, we intend to conduct further analysis to determine whether, in the absence of diagnosis code modifications, there are alternative configurations to classify cases in DRGs 121-123 that better predict resource use among patients. We would also urge hospitals performing cardiac catheterizations on AMI patients to code those catheterizations even though they do not affect the DRG assignment at this time, as more complete data on procedures performed is of value in assessing the appropriateness of classification changes.

### 4. Adding a CC to DRG 124

We proposed to add diagnosis code 428.9, Heart Failure, unspecified, to the diagnoses included in DRG 124 (Circulatory Disorders except AMI with Cardiac Catheterization and Complex Diagnosis). We received no negative comments and one favorable comment

in support of this proposal. We are adopting it without change.

### 5. Pacemakers

We proposed to continue to classify single and dual chamber pacemakers in the existing pacemaker DRGs, 115 through 118.

Comment: One commenter wrote expressing displeasure with our failure to reclassify pacemaker cases based on the type of device. The commenter encouraged further study in this area so that hospitals will not continue to be underpaid for dual chamber devices in the future.

Response: As we explained in the proposed notice, the pacemaker classification issue is more complicated than identifying and recognizing price differences between dual chamber and single chamber devices. Moreover, new ICD-9-CM codes, which will permit us to distinguish among types of pacemaker devices, have just been adopted for use effective in October 1987. These codes may be found in Table II of this notice. We note that ProPAC, as well as several manufacturers of pacemaker devices, support our decision to conduct further study on this issue. We will report our findings as they become available.

Although we proposed no classification changes for cardiac pacemaker cases, we note that the adoption of new ICD-9-CM codes for cardiac pacemaker procedures requires modification of the Grouper logic in order to ensure consistent treatment of like cases using both the old codes and the new codes. A summary of the revised Grouper logic for classifying pacemaker cases appears as Table III in Section VI of this notice.

### 6. Defibrillators

We proposed the following interim measures for reclassification of automatic implantable cardioverter defibrillator (AICD) cases: to continue to assign AICD total system implants with cardiac catheterization to DRG 104 (Cardiac Valve Procedure with Pump and With Cardiac Catheterization); to assign AICD total system implant cases without cardiac catheterization to DRG 105 (Cardiac Valve Procedure with Pump and Without Cardiac Catheterization); and to assign procedure codes 37.95 through 37.98, Implant or replacement of cardiodefibrillator leads or generator, to DRG 120 (Other Circulatory System OR Procedures), and remove them from DRG 117 (Cardiac Pacemaker Revision Except Device Replacement).

Comment: Two commenters expressed concern with our proposed classification of defibrillator cases. The commenters expressed concern that ICD-9-CM coding conventions required clarification so that electrophysiological testing performed in association with the implant is reported. One commenter also believes that reclassification of defibrillator replacements and/or defibrillator lead implants from DRG 117 to DRG 120 would continue to result in underpayment for the procedure. The commenter requested reclassification to an unspecified higher weighted DRG.

Response: We admit that there is not a distinct ICD-9-CM code for electrophysiological testing. Currently, this is captured under code 37.29, Other diagnostic procedures on the heart and pericardium. The ICD-9-CM Coordination and Maintenance Committee addressed this issue at their July 1987 meeting. During the coming year it is anticipated that a new code will be proposed for electrophysiological testing. If approved, the code(s) would become effective October 1, 1988. In addition, hospitals should report cardiac catheterization on the Medicare billing form whenever performed. Use of the catheterization code affects DRG classification in several DRGs even though it is not considered an operating room procedure.

With regard to the classification of replacement and/or insertion of AICD leads and devices alone, we continue to believe DRG 120 is appropriate for the time being. Based on the limited Medicare data available at this time, we find that average charges for AICD procedures other than total system implants reasonably approximate the average charge for DRG 120. Until more claims data are available, we consider it more appropriate to temporarily classify the AICD procedures, other than total system insertions, in DRG 120. Once the technology is stabilized and we have a larger data base for evaluative purposes, we will consider further reclassification of both total system AICD implants and other defibrillator procedures. We note also that ProPAC's comments include support for assignment of these procedures to DRG 120.

Comment: One commenter noted our commitment, published in the September 3, 1986, PPS notice to reevaluate complex aortic aneurysm repairs. The commenter expressed disappointment that the results of this re-evaluation were not contained in the May 19, 1987 proposed notice and requested that the results of the evaluation be included in this notice.

Response: The new ICD-9-CM procedure codes that clearly identify thoraco-abdominal aortic aneurysm repair just became effective October 1, 1986. Because of the lag between patient

discharges and inclusion of the discharge data in our central office MEDPAR file, we have limited data from FY 1987.

We believe it is premature to conduct a re-evaluation on complex aortic aneurysm repairs at this time due to the scarcity of data. Any conclusions based on such limited data would of necessity be considered interim, as with cochlear implants and cardiac defibrillators. Since we have already instituted an interim reclassification of the procedure. we do not believe it is a prudent use of resources to engage in a rushed analysis based on incomplete data, only to repeat the analysis once more information becomes available. Consequently, we have nothing to report on this procedure at this time.

We do intend, however, to re-evaluate the classification of complex aortic aneurysm repairs once adequate data are available. We will report on our findings in the first DRG classification notice to be published after our review is complete.

D. MDC 6: Diseases and Disorders of the Digestive System

1. Reassignment of Ulcer Diagnosis Code

We proposed to remove diagnosis code 531.70, Chronic gastric ulcer without mention of hemorrhage, perforation or obstruction, from DRG 176 (Complicated Peptic Ulcer) and reassign it to DRGs 177 (Uncomplicated Peptic Ulcer with CC) and 178 (Uncomplicated Peptic Ulcer without CC). We did not receive any comments on this proposal, so it will be adopted without change.

2. No Change in the Classification of Bowel Procedures

We stated that national data indicated no significant classification problems in the major bowel procedure DRGs 148 and 149 (Major Small and Large Bowel Procedures). Thus, we proposed not to amend the current classification of bowel procedures. We did not receive any comments on our statement, so the current classification of bowel procedures will remain.

E. MDC 8: Diseases and Disorders of the Musculoskeletal System and Connective Tissue

### 1. Hamartoma

We proposed to remove diagnosis code 759.6, Other hamartoses, not elsewhere classified (NEC), from MDC 17, Myeloprofiferative and Poorly Differentiated Neoplasms, and to add it to MDC 8. We also proposed adding procedure code 32.29, Other local excision or destruction of lesion or tissue of lung to DRGs 233 and 234 Other Musculoskeletal and Connective Tissue Procedures), in order to avoid inappropriate assignment to DRG 468.

Comment: We received conflicting comments concerning our proposal to move diagnosis code 759.6, Other hamartoses, NEC, from MDC 17 to MDC 8. One commenter approved of the proposed change but, noting that hamartoses occur at sites other than the lung, recommended that additional procedure codes be added to DRGs 233 and 234 to prevent assignment to DRG 468 of excisions of hamartoses at other sites. Another commenter expressed concern that hamartoses not be removed from MDC 17, stating that a hamartoma is a tumor-like mass that may undergo neoplastic transformation. Therefore, the commenter believes assignment to MDC 17 is appropriate.

Response: In light of the conflicting comments on this subject we have decided to postpone implementation of this proposed change for FY 1988. We will study the issue in greater detail during the upcoming evaluative period and will report on our findings in next

year's proposed notice.

### 2. Certain femur procedures

We proposed to reassign procedure code 79.95, Unspecified operation on femur injury from DRGs 218, 219, and 220 (Lower extremity and humerus procedure except hip, foot, femur). We received no negative comments and one favorable comment in support of this proposal, so it will be adopted without change.

F. MDC 9: Diseases and Disorders of the Skin, Subcutaneous Tissue and Breast

### 1. Cellulitis of finger, toe and digit

We proposed to add the following diagnoses to the list of principal diagnoses that may result in assignment to DRG 263 (Skin Graft and/or Debridement For Skin Ulcer or Cellulitis with CC) and DRG 264 (Skin Graft and/ or Debridement For Skin Ulcer or Cellulitis without CC).

681.00 Cellulitis and abcess of finger, unspecified

681.01 Felon

681.02 Onychia and paronychia of finger

681.10 Cellulitis and abscess of toe, unspecified

681.11 Onychia and paronychia of toe 681.9 Cellulitis and abscess of unspecified digit

We received no negative comments on this proposal, so it will be adopted without change.

2. Infected abrasion or friction burn of face, neck, and scalp

We proposed to remove diagnosis code 910.1, Abrasion or friction burn of face, neck, and scalp except eye, infected, from DRGs 280-282 (Trauma to the Skin, Subcutaneous tissue, and Breast), and assign the code to DRGs 277-279 (Cellulitis) along with other diagnoses for infection.

We received no negative comments on this proposal, so it will be adopted

without change.

G. MDC 11: Diseases and Disorders of the Kidney and Urinary Tract

We proposed to remove diagnosis code 581.9, Nephrotic syndrome with unspecified pathological lesion in kidney, from DRGs 325-327 (Kidney and Urinary Tract Signs and Symptoms) and place it in DRGs 331-333 (Other Kidney and Urinary Tract Diagnoses). We did not receive any negative comments on this proposal, so it will be adopted without change.

H. MDC 18: Infections and Parasitic Diseases (Systemic and Unspecified Sites)

We proposed to remove diagnosis code 785.59, Shock without mention of trauma, NEC, from MDC 5 and reassign the condition to MDC 18, with nonsurgical cases assigned to DRGs 416 and 417 (Septicemia). We did not receive any comments on this proposal, so it is adopted without change.

I. MDC 20: Alcohol/Drug Use and Alcohol/Drug Induced Organic Mental Disorders

### Refinement of alcohol/drug DRGs

In accordance with our September 3, 1986 final rule on changes to the inpatient hospital prospective payment system and FY 1987 rates (51 FR 31454). the exclusion of alcohol/drug treatment hospitals and units was extended through cost reporting periods beginning before October 1, 1987. The extension was intended to permit completion of analyses of a record reabstraction study conducted by the Alcohol, Drug Abuse, and Mental Health Administration (ADAMHA) in concert with the National Institute of Mental Health, the National Institute on Drug Abuse, the National Institute on Drug Abuse, the National Institute on Alcohol Abuse and Alcoholism, the Office of the Assistant Secretary for Planning and Evaluation,

The record reabstraction study was designed to examine the predictive value of variables not currently included in the DRG logic for MDC 20, such as patient age, disability status, CC and

polysubstance use, on patient resource use. Based on the analyses and recommendations of ADAMHA, HCFA tested the effects of reconfiguring the alcohol/drug DRGs along the lines suggested by ADAMHA by analyzing the FY 1985 and FY 1986 MEDPAR records for all Medicare discharges in MDC 20 (the ADAMHA reabstraction study was based on a stratified random sample of FY 1984 discharges). Based on our analyses and those of ADAMHA. we proposed to reconfigure the alcohol/ drug DRGs as follows:

· Cases in the current DRGs 434 and 435 would be combined and then re-split based on the presence or absence of non-MDC 20 CC. The proposed DRG 434 would be Alcohol/Drug Abuse or Dependence, Detoxification or Other Symptomatic Treatment, with CC and the proposed DRG 435 would be Alcohol/Drug Abuse or Dependence, Detoxification or Other Symptomatic Treatment, without CC.

· We proposed that cases with either a principal or secondary diagnosis (rather than only a principal diagnosis, as currently) of alcohol or drug dependence would be assigned to the revised DRG 436, if rehabilitation therapy (diagnosis code V57.89) was furnished, and to the revised DRG 437 if both rehabilitation therapy and detoxification (procedure code 94.25) were provided. In addition, we proposed that certain diagnoses that are not currently assigned to DRGs 436 and 437 be permitted to group to the revised DRGs 436 and 437 if it is the physician's judgment that a patient with a principal or secondary diagnosis of one or more of these conditions may benefit from rehabilitation and such rehabilitation is furnished and reported on the bill submitted. These additional diagnoses to be treated as dependence diagnoses were alcohol amnestic syndrome (291.1). other alcoholic dementia (291.2), alcohol withdrawal hallucinosis (291.3), other specified alcoholic psychosis (291.8). and unspecified alcoholic psychosis

The revised logic for DRGs 436 and 437 would move cases involving the above diagnoses from DRG 434 (to which codes 291.1 and 291.2 are currently assigned) and from DRG 435 (to which codes 291.3, 291.8 and 291.9 are currently assigned) as long as rehabilitation therapy was furnished. The effect of recognizing secondary as well as principal diagnoses of alcohol or drug dependence in the proposed revisions of DRGs 436 and 437 was to remove cases from DRG 434 when rehabilitation therapy was furnished to a patient with a principal diagnosis of

alcohol or drug abuse, as long as the record also contained evidence of alcohol or drug dependence as a secondary diagnosis.

No change was proposed to the logic for DRG 433, except that it be renamed Alcohol/Drug Abuse or Dependence, Left Against Medical Advice.

We received several comments on these proposed DRG revisions. We also received numerous comments on the scheduled end of the exclusion of alcohol/drug hospitals and units, the absence of an impact analysis on the effect of inclusion of alcohol/drug hospitals and units in the prospective payment system, the proposed recalibrated weights, and outlier thresholds for the alcohol/drug DRGs. They will be addressed separately in our final rule of changes to the inpatient hospital prospective payment system and FY 1988 rates, published elsewhere in this issue of the Federal Register.

Comment: Several associations representing hospitals, physicians and nurses commented favorably on the proposed restructuring of DRGs 434-437. indicating that the proposed DRGs for alcohol/drug diagnoses represented an improvement over the current classifications. The National Association of Addiction Treatment Providers (NAATP), while expressing major concerns regarding the recalibrated DRG weights, the scheduled end of the exclusion of alcohol/drug hospitals and units and the absence of an impact analysis, wrote: "We sincerely applaud the Department's efforts and diligence represented by the various analyses which have been conducted and served as the basis for the current proposed reclassification of the alcohol and drug abuse DRGs. We think that the proposed reclassification has a solid analytic foundation and fairly represents a method to classify alcoholism and drug abuse/dependency admissions for the Medicare beneficiary population."

Response: We are encouraged by the favorable response to the proposed reconfiguration of the alcohol/drug DRGs. We note that while many individual alcohol/drug hospitals and units were critical of the scheduled end of their exclusion from the prospective payment system, their criticism related to the absence of an impact analysis and the use of cases from short-stay hospitals as well as excluded alcohol/ drug hospitals and units to establish the proposed weighting factors, rather than to the proposed classification of alcohol/drug cases among the MDC 20 DRGs. We believe this is further evidence that the proposed DRG classifications do represent an

improvement over the existing alcohol/drug DRGs.

Comment: One commenter expressed concern that the June 10 notice provided only general descriptions of the ADAMHA and HCFA studies on which the proposed modifications were based but did not display the study results. This commenter also indicated that the study files had not been made available

to the public.

Response: The ADAMHA study was designed to assess and test hypotheses regarding alternate structuring of the MDC 20 DRGs. The findings of this reabstract study were published in the June 10 proposed notice as supporting evidence for the proposed DRG reconfigurations. Sufficient information was provided to document the impact of those variables found to influence resource consumption in the alcohol/ drug DRGs. Our description of the analyses undertaken identifies the data sources used and provides sufficient information to enable interested parties to conduct similar analyses, should they wish to do so. The availability of MEDPAR data, on which most of HCFA's analyses are based, was the subject of a Federal Register notice published July 2, 1985 (50 FR 27407). The ADAMHA reabstraction study database, consisting of data reabstracted from the medical record merged with MEDPAR data, contains personally identifiable data which under the provisions of the Privacy Act, may not be released. ADAMHA is in the process of removing personal identifiers and will make the data available to the public at the earliest possible date. Preparation of this data tape has involved careful screening of all variables to assure that no information identifying specific patients or providers is included. All provider-specific variables that might possibly identify a given provider were omitted, in support of ADAMHA's assurance of confidentiality with respect to the identities of providers participating in the study and their patients whose records were abstracted. Also, as noted earlier in this document, a contact person's name and telephone number are provided for those seeking additional information on the changes. Finally, all materials used and analyses conducted are subject to release upon request, including requests under the Freedom of Information Act. We believe that all of these procedures facilitate open, public debate and permit readers sufficient information in sufficient detail, to evaluate the proposed changes.

Comment: One commenter believed that the alcohol/drug DRGs should differentiate between patients with dual diagnoses, such as cirrhosis, depression, or other medical or psychiatric diagnoses, and those without. Other commenters thought presence or absence of CC should be incorporated into the revised DRGs 436 and 437.

Response: We evaluated the impact of presence or absence of any CC on all alcohol/drug DRGs and found that most patients with a principal diagnosis in MDC 20 also had a CC reported. Because most alcohol/drug diagnoses are on the master CC list and many patients in MDC 20 have multiple alcohol/drug diagnoses, such as both dependence and abuse, coded, this distinction was not associated with significant differences in resource use. After revising the CC list for MDC 20 to exclude diagnoses in MDC 20, we found that presence or absence of non MDC-CC was associated with significant differences in resource use among patients not undergoing rehabilitation.

Based on FY 1986 MEDPAR records. patients in DRGs 434 and 435 with non-MDC 20 CC had mean charges 40 percent and 45 percent greater, respectively, than patients without CC. Moreover, the homogeneity of the resulting groups, as measured by a weighted average of the coefficients of variation, improved by 3 percent for DRG 434 and by 8 percent for DRG 435. Because there were no statistical differences between charges for patients in DRGs 434 and 435 when presence or absence of CC was held constant, we proposed to combine all cases from DRGs 434 and 435 and then re-split them into two new DRGs, distinguished by presence or absence of non-MDC 20 CC. Accordingly, we believe our proposed revision of DRGs 434 and 435 does take into account differences between patients with multiple diagnoses and those without, as recommended by the first commenter cited.

When we performed similar analyses on cases in DRGs 436 and 437, we found that the resource differences between patients with CC and those without dropped to 17 and 10 percent, respectively. Moreover, the homogeneity of the two resulting pairs improved by only one-half of one percent. We infer from this minimal improvement in homogeneity that there are other factors contributing nearly as much to patient differences in resource use as the presence or absence of CC. As we indicated in the proposed notice, the average resource differential in over 100 pairs of DRGs, when split on the basis of CC, averages 60 percent for all pairs and is less than 20 percent for only two pairs. In light of the more modest resource differences associated with CC

in DRGs 436 and 437, combined with the minimal improvements in homogeneity of the resulting groups, we do not believe it is appropriate at this time to split DRGs 436 and 437 on the basis of presence or absence of CC. Therefore, we have not adopted the commenters' recommendations regarding DRGs 436 and 437.

Comment: One commenter observed that resource differences associated with multiple addictions should be reflected in the revised MDC 20 DRGs. Another commenter, while acknowledging that the proposed alcohol/drug DRGs are an improvement over the current DRGs in MDC 20, questioned the conclusions implied by a lack of differentiation in the DRGs between the types of substances abused

between the types of substances abused. Response: At the present time, cases of drug abuse represent less than fifteen percent of all Medicare cases in MDC 20. However, ADAMHA's study sample was designed so as to reabstract all Medicare alcohol/drug cases from each provider in the sample; it was hypothesized that differential resource intensity associated with type or combination of substances used would emerge at the provider level.

In an effort to evaluate the effects of multiple substance abuse ADAMHA examined cases involving alcohol only, single drug use only, alcohol/drug combinations, and multiple drug combinations. Based on the limited number of Medicare cases involving drug use other than alcohol, ADAMHA did not consider it appropriate to distinguish the DRGs by type of drug used or between alcohol and drug cases, nor did it find systematic and consistent results when it distinguished cases involving single substance abuse from those involving polysubstance abuse. On the basis of these analyses, ADAMHA recommended no distinction in the DRGs between alcohol and drug dependence or between single substance and multiple substance

Comment: One commenter noted that the proposed notice referenced the use of code V57.81 for rehabilitation rather than code V57.89.

Response: We agree with the commenter that the proposed notice contained a typographical error. The correct code for alcohol/drug-related rehabilitation is V57.89. We regret any confusion this error may have caused our readers.

Comment: One commenter raised questions about the validity of the ADAMHA study in light of the sample size and the use of only the face sheet and discharge summary as the basis for reabstraction.

This commenter observed that acceptable coding practices require use of the entire medical record, especially to identify CC.

Response: ADAMHA consulted extensively with industry representatives, HCFA and the Assistant Secretary for Planning and Evaluation regarding the design, implementation and analyses conducted during their MDC 20 study. The ADAMHA reabstraction study was based on a stratified random sample of 849 hospitals and units with some 15,000 Medicare discharges in the alcohol/drug DRGs. These discharges represented 28 percent of all alcohol/drug cases in FY 1984. Providers were stratified by type of hospital or unit and its PPS status (excluded hospital, excluded alcohol unit, included short-stay hospital, etc.). Within this sampling frame, sampled providers were asked to submit records for all of their MDC 20 discharges for FY 1984. The provider response rate approached 90 percent, and 80 percent of the requested records were obtained. Even after adjusting for response rate, the reabstracted records comprise 22 percent of Medicare alcohol/drug cases in FY 1984. We believe this sampling frame yielded an adequate database for analysis from the standpoint of both size and representativeness of the selected providers.

As to the validity of using only the face sheet and discharge summary as the basis for reabstraction, it is true that coders are encouraged to review the entire medical record in order to capture any diagnoses that might have been omitted by the physician who prepared the discharge summary. However, ADAMHA conducted a pilot test to compare the availability and reliability of variables of interest from the discharge summary with their availability from the full medical record. The pilot test confirmed that the face sheet and the discharge summary were in fact valid and reliable sources of the necessary information and demonstrated that their use would facilitate efficiency of the data collection effort without sacrificing accuracy. In addition, the study methodology specifically provided for call-back consultation with hospital medical records personnel and involved referring back to the full medical record whenever there was any doubt about the variables of interest being reabstracted from the face sheet and the discharge summary.

Coders working on the reabstraction study were instructed specifically to code coexisting conditions. The data collection instrument permitted coding of up to seven diagnoses and five procedures, in contrast to the maximum of five diagnoses and three procedures that can be reported on the Medicare billing form. The ADAMHA study paid particularly close attention to accurate coding of the incidence of rehabilitation.

We believe the procedures followed by ADAMHA in designing and conducting this study were sufficient to ensure completeness, quality and consistency of the data collected and the reliability of inferences drawn with respect to analyses of alternative configurations of the alcohol/drug DRGs.

Finally, we reiterate that the final analysis on many of the alternative configurations of the MDC 20 DRGs were ultimately tested on more recent and more complete Medicare data on discharges from the alcohol/drug DRGs—more than 40,000 in FY 1985 and more than 30,000 cases in FY 1986 all discharge data received in HCFA central office through September 1986.

Comment: One commenter observed that because the current MDC 20 DRG assignments are not affected by the presence of CC, it is likely that such conditions were under-reported in the ADAMHA study and in FY 1986

Response: As noted previously, ADAMHA specifically collected comprehensive data on secondary diagnoses, using a data collection instrument designed to collect up to seven diagnoses rather than five as are reported on the hospital billing form. In the FY 1985 and FY 1986 MEDPAR study files, the percentage of patients in the alcohol/drug DRGs with non-MDC 20 CC on their Medicare hospital bills was comparable to the percentage of patients in other DRGs with CC. We wish to emphasize that we require reporting of CC on all Medicare cases, even in DRGs not affected by the presence or absence of CC. We encourage hospitals to code diagnostic and procedural information as completely and accurately as possible on all Medicare claims. Complete and accurate medical record coding not only ensures the accuracy of DRG assignment; it also enriches the Medicare program data bases used for our ongoing monitoring and evaluation of the prospective payment system, including DRG classification changes.

Based on the foregoing discussion of comments on the proposed revisions of the MDC 20 DRGs, we are adopting our proposed reconfiguration of DRGs 434–437 without change.

#### IV. Proposed Changes and Comments Affecting Multiple MDCS

#### A. Elimination of Age Over 69

In the May 19, 1987 proposed notice, we proposed to eliminate age over 69 as a criterion for DRG classification in all of the pairs of DRGs in which age over 69 and/or CC was a factor. We also proposed to continue to consider age under 18 in those pediatric DRGs that already have been established and to continue to consider age 35 in the diabetes DRGs (294 and 295).

Comment: Several rural hospitals wrote expressing dissatisfaction with our proposed elimination of age over 69 as a criterion for DRG classification. These commenters believed that the proposed change would result in a significant loss of revenue to rural hospitals. One commenter observed that small rural hospitals were not in a position to look for and code esoteric CC in a patient population comprised disproportionately of elderly Medicare beneficiaries.

Another commenter, a hospital, indicated that it had evaluated the impact of this proposal on 75 Medicare patients age 70 or older. It noted that 59 of the patients would be unaffected by the change but that it would experience a substantial drop in revenue for the

remaining 16 cases.

Response: We believe these comments reflect legitimate concern mixed with some misunderstanding of the effects of the proposed change. About 70 percent of Medicare beneficiaries hospitalized each year are age 70 or older. About 60 percent of Medicare beneficiaries under age 70 are reported to have CC. As a result, the current split in the paired DRGs based on age over 69 or presence of CC results in about 88 percent of all Medicare cases in the paired DRGs being assigned to the DRGs involving age over 69 and/ or CC. The fact that such a substantial majority of Medicare beneficiaries are grouped into these DRGs has the effect of masking significant differences in resource use among patients within the DRG pair because the DRG for older and/or complicated cases includes a mix of extremely sick patients who are very resource-intensive and patients who present no complications and are not very different from younger patients in their average resource use.

When age greater than 69 is eliminated from consideration, the differences between the resulting DRGs—those with and those without CC—generally widens and the variance within the resulting DRGs is less than in the former. In most pairs, the weighting factor for the DRG with CC rises and the

weighting factor for the DRG without CC declines relative to the current weighting factors. (It is important to note that these are the combined results of both reclassification and recalibration; we have not isolated the effects of using FY 1986 data to recalibrate the weights separately from the DRG classification changes.) Accordingly, in the example described by one commenter, if the weighting factor and, hence, payment for 16 Medicare cases was projected to decline, the weighting factor for the 59 'unaffected" cases would rise. We believe this outcome was not understood by the commenters.

Even after elimination of age over 69 as a criterion in DRG assignment, some 60-65 percent of Medicare beneficiaries will continue to be grouped to the complicated DRGs within the pairs. Data on the distribution of complications by age reveals that there is a steady rise in the incidence of CC as age increases, from 58 percent among 65-year-olds to 71 percent for patients age 85 and older. To the extent that rural hospitals treat disproportionately more older patients among the Medicare population, we would expect to see disproportionately more of their cases assigned to the DRGs with CC. As to the allegation that rural hospitals are not in a position to identify esoteric CC, we note that there are about 2,700 diagnoses on the master CC list, the majority of which are not in the least esoteric but rather are diagnoses which, when principal, are assigned to some of the most frequently occurring DRGs in

the Medicare population.

We should also point out that the data demonstrate that the presence of CC contributes much more significantly to the variation in case costs than does age over 69. After controlling for CC, patients over age 69 are, on average, four percent more expensive than patients under age 70, whereas, regardless of age, patients with complications are about 30 percent more expensive than patients without complications. This finding emerges in analyses conducted by ProPAC and HCFA and is consistent across three consecutive years of Medicare data (FYs 1984, 1985, and 1986). We do not believe it is within the overall interest of the general public to ignore those results.

Supporters of maintaining age over 69 in the DRG classification scheme argue that older patients generally require more resources than younger counterparts. Despite the fact that a few patients age 70 and older without CC are as expensive to treat as patients with CC, the data show that this is not generally the case. Consequently, the

commingling of all aged patients with patients presenting CC reduces the mean for those DRGs and, thus, tends to result in underpayment for more complicated cases.

Since the DRG classification system is based on averages, there will always be cases with costs both above and below the payment level. Given that there is a fixed data base from which the weighting factors are computed. classification changes generally result in increased payment for one kind of case at the expense of decreased payment for another kind of case. Our intent is to refine the classification systems to be as equitable as possible. Thus, even though hospitals will experience a decrease in revenue for uncomplicated older patients, they will experience an increase in revenue for complicated cases. The net effect on an individual hospital is dependent upon its mix of patients.

Finally, we do not expect hospitals to frequently encounter situations in which an older patient who requires extremely intensive resources does not have any CC. The classic example cited of a 90-year-old patient utilizing a high level of nursing services without presenting any CC, simply does not occur frequently in reality. There is a high correlation between the intensity of services and the presence of CC. Consequently, we believe the elimination of age over 69 from the classification structure results in payments that better reflect resource use.

Comment: One commenter stated that both HCFA s and ProPAC's analyses are limited by their failure to consider alternative age splits that may be more meaningful in the Medicare population. Another commenter recommended that the complicated versus uncomplicated DRGs be further split on basis of age

Response: Although not specifically addressed in the proposed notice, we evaluated alternative age splits in our research on this issue. We found that alternative age breaks, in combination with CC, produced no overall improvement in the homogeneity of the DRGs. In addition, neither we nor ProPAC found that age by itself explained as much variation in charges as did the presence of CC. Regarding the recommendation that the revised DRGs be further split by age, thus doubling the number of DRGs in pairs, ProPAC found that patients over age 69 without CC were, on average, 4 percent more expensive than patients under age 70 without CC. This small difference does not justify duplicating DRGs on the basis of age.

Comment: Several commenters encouraged HCFA to split all DRGs based on presence of CC.

Response: We agree that it is necessary to investigate whether the presence of a CC significantly affects resource consumption in DRGs other than those that are currently split on that basis. We will be evaluating that issue and will report on our findings in the future.

Comment: One commenter voiced dissatisfaction with the elimination of age over 69 and argued that all DRGs must be reweighted in order to properly

reimburse hospitals.

Response: The weighting factors for all the DRGs have been recalibrated for FY 1988 and annually thereafter. As described in our proposed notice, the weighting factors are based on standardized charges for all Medicare discharges in FY 1986 for which data have been received through June 1987. Those cases are grouped in accordance with the logic that will determine DRG assignment in FY 1988, including all of the classification changes finalized in this notice. The average case weight before and after DRG reclassification and recalibration of the weights is held constant to ensure that reclassification and recalibration neither increase nor decrease projected Medicare outlays for the same set of cases. Thus, the elimination of age over 69 as a factor in DRG assignment, as well as all other DRG classification changes, has been taken into account in computing the revised DRG weights. The DRG weights listed in table 5 of the addenda to both the June 10 proposed rule and the final rule published elsewhere in this issue reflect this recalibration.

Comment: Two commenters noted our statement that the elimination of age over 69 resulted in some DRGs being more internally heterogeneous. One commenter, briefly summarizing some of its own research, argued that the relationship between age and charges depends on the DRG weight. The commenters encouraged additional evaluation of the age proposal, possibly resulting in the splitting of certain DRGs

by age and CC.

Response: When initially evaluating the appropriateness of eliminating age over 69, we considered the idea of making individual determinations as to whether to retain age in the classification of each pair of DRGs split on the basis of age and/or CC. However, we rejected this alternative primarily because we believe that annual evaluation, potentially causing some DRGs to consider age one year and not the following, would be extremely confusing and burdensome to hospitals

as well as to HCFA and other DRG users. We believe it is preferable to sacrifice a minimal amount of internal homogeneity in a few DRG pairs than to present users with the uncertainty of not knowing the DRGs that would be split on age each year. Moreover, we do not believe such an annual re evaluation of each of the paired DRGs is a prudent use of resources, nor do we believe that it would be equitable to make a permanent decision to retain age over 69 for a few DRGs based on a single year's findings.

Comment: Several other commenters objected to the elimination of age over 69 as a factor in DRG classification. Two of the commenters stated that the proposal would adversely affect their hospitals because older patients require a greater intensity of nursing services that would no longer be reflected in payment. Another commenter noted that aged patients utilize a disproportionate share of non-revenue-producing hospital resources, such as social services, which should be considered.

Response: We continue to believe that age over 69 should be eliminated from the DRG classification system. The data indicate that the presence of CC has much greater power than either age by itself or age in combination with CC in explaining variations of resource use among patients, as measured by standardized charges. While we admit that the nursing needs of a healthy 90year-old may be somewhat more than a healthy 65-year-old, the nursing needs of either a 65-year-old or a 90-year-old with CC are likely to exceed those of either patient without CC. Furthermore, we suspect that the vast majority of patients who require a greater intensity of nursing services do, in fact, present CC. We should point out that by not commingling relatively healthier, older patients with the patients who present CC, the average charge for the complicated cases increases. Consequently, the proposed revision in the classification system results in payment that more accurately reflects the additional services furnished.

With regard to non-revenue-producing hospital resources, such as social services, we must point out that the Medicare program, or any other payer of which we are aware, has never recognized differential utilization of such services. Rather, the cost of non-revenue-producing services are allocated to revenue producing services. Consequently, we have no means of determining the relationship between such services and age as opposed to CC.

B. Addition of Drug Diagnoses to CC List

In our May 19 notice we proposed adding the following diagnoses to the list of CC.

304.20 Cocaine dependence, unspecified

304.21 Cocaine dependence, continuous

304.22 Cocaine dependence, episodic 304.50 Hallucinogen dependence, unspecified

304.51 Hallucinogen dependence, continuous

304.52 Hallucinogen dependence, episodic

304.60 Other specified drug dependence, not elsewhere classified, unspecified

304.61 Other specified drug dependence, not elsewhere classified, continuous

304.62 Other specified drug dependence, not elsewhere classified, episodic

305.30 Hallucinogen abuse, unspecified 305.31 Hallucinogen abuse, continuous

305.32 Hallucinogen abuse, episodic 305.60 Cocaine abuse, unspecified

305.61 Cocaine abuse, unspecified

305.62 Cocaine abuse, episodic

305.90 Other, mixed, or unspecified drug abuse

305.91 Other, mixed, or unspecified drug abuse, continuous

305.92 Other, mixed, or unspecified drug abuse, episodic

C. Refinement of Complications and Comorbidities Listing

In the May 19 notice we proposed to modify the Grouper logic so that certain diagnoses generally included on the list of CC would not be considered a valid CC in combination with a particular principal diagnosis.

Comment: One commenter noted that many principal diagnoses listed fall into MDCs that are not split on the basis of CC, such as mental disorders. The commenter believes it is pointless to complicate the task of refining the CC list by including these codes.

Response: We had considered limiting the CC refinement when we initially undertook this task. However, given the ongoing process of DRC refinement, future work of evaluating all DRGs for additional CC splits, and general research in classification methodologies, we believed it best to design the CC listing to be as all-encompassing as possible. Thus, if future DRG changes require CC splits for additional DRGs, the CC exclusion listing will be able to handle the change adequately without

massive modification of the CC exclusion list.

We regret any confusion this process may have created for some reviewers. However, we continue to believe it is prudent to remain prepared for as many future alternatives as possible in developing such a major revision to the software space utilized by the Grouper program.

Comment: Two commenters noted typographical errors in the CC exclusion

listing.

Response: Although we attempted to proofread the copy of the CC exclusion listing carefully, a few errors were identified during the review of over 350,000 codes. The following changes should be made to the listing.

1. On page 1, principal diagnosis code 006.4 should read 006.5. Then, principal diagnosis code 006.4 should be added with code 513.0 excluded as a CC.

- 2. On page 515, principal diagnosis codes 807.00, 807.01, 807.02, and 807.03 were omitted inadvertently. They should be added with the same list of exclusions as under principal diagnosis code 807.05.
- 3. On page 687, under principal diagnosis code 868.12. all exluded codes listed should be deleted and replaced with only code 868.12.
- 4. On page 711, under principal diagnosis code 998.1, excluded code 981 should read 998.1.
- 5. The following codes are deleted from the list of excluded diagnoses and will continue to be treated as CC:
- (a) On page 56, delete code 723.4 from exclusions listed under principal diagnosis code 054.40,
- (b) On page 119, delete code 424.0 from exclusions listed under principal diagnosis code 395.0
- (c) On page 151, delete codes 607.1, 607.2, and 607.3 from exclusions listed under principal diagnosis code 599.7.
- (d) On page 199, delete codes 614.0, 614.3, 614.5, 615.0, 616.3, 616.4, and 620.7 from exclusions listed under principal diagnosis code 753.9,
- (e) On page 199, delete all codes that do not begin with a "7" from exclusions listed under principal diagnosis code 759.8.
- (f) On page 515, delete code 070.5 from exclusions listed under principal diagnosis code 807.04,
- (g) On page 696, delete all codes from exclusions listed under principal diagnosis codes 901.83, 901.89, and 901.9.
- (h) On pages 121–122, delete code 427.5 from exclusions listed under principal diagnosis codes 426.0, 426.10, 426.11, 426.12, 426.13, 426.2, 426.3, 426.4, 426.50, 426.51, 426.52, 426.53, 426.54, 426.6, 426.7, 426.81, 426.89, and 426.9.

6. Delete from the exclusions list, wherever it appears, any diagnosis code that is not already on the master list of CC (as modified by this notice). For example, on page 1, the first exclusion listed under principal diagnosis 003.1 is code 020.2. Since code 020.2 is not already a CC, it need not be excluded and should be deleted whenever it appears on the CC exclusions list.

Comment: One comment noted that some codes remain valid CC for themselves while other codes exclude themselves as valid CC. The commenter believes the revised CC exclusions for coding should be consistent.

Response: Whether or not a particular code can be a CC for itself was based on clinical judgment of the medical consultants developing the CC listing. A very few codes were not excluded from themselves because they may signify the bilateral occurrence of particular

condition.

Other situations in which a code is not excluded from itself represent diagnoses that are not on the overall list of CC to begin with and thus need not be excluded from the CC list for any principal diagnosis. Such is the case in the example cited by one commenter of code 349.89, Kline Levin Syndrome. Since code 349.89 is not on the master CC list, it is entirely unnecessary to exclude it from the CC list for itself or any other principal diagnosis. In addition, as noted above, we have deleted from the list of exclusions any diagnosis code that was not already on the master CC list in an effort to eliminate any confusion regarding whether a particular diagnosis is ever a

Comment: One commenter, anticipating that it would be impossible for hospitals and other organizations to incorporate the CC refinement into revised software by October 1, 1987, expressed concern that implementation of the CC refinements would disrupt billing operations. Given the elimination of periodic interim payments (PIP) effective July 1, 1987, this commenter recommended that HCFA make only necessary changes to data processing systems to minimize delays in the billing process.

Response: Hospitals are not required to make any changes in the information they furnish in order to have Medicare bills processed after the implementation of the CC refinement. Hospitals are free to report diagnoses on their Medicare billing form just as they have been doing since 1983. The Medicare FIs assign the DRG based on diagnosis and procedure codes reported. The DRG reported by hospitals is not considered by the FI in making payment. Consequently,

hospitals should continue to submit their Medicare bills timely despite the fact that their internal Grouper software may not be revised by the beginning of the fiscal period.

Comment: One commenter noted that both the original CC listing and the proposed refinement to the listing were based on clinical judgments. The commenter believes that the CC list and exclusions need to be based on a more empirical foundation.

Response: We agree that empirical findings combined with clinical judgment should be considered in any broad refinement of the CC listing. We currently have let a research grant in this regard. We anticipate further improvements in the CC listing, once the research is completed.

In the meantime, we do not believe the absence of empirical findings should delay implementation of the currently proposed refinement to the CC listing. We believe that the proposed exclusions to the CC listing adhere closely to the principles that guided this endeavor, as described in the proposed notice. As we have stated above, we will continue to monitor the impact of the refinement and make further revisions based on empirical findings as they become available.

Comment: Several commenters wrote expressing general concern with the detail of the CC refinement. Two of the commenters were not confident that the listing was complete and urged us to ensure that all diagnoses that should not be considered as CC for a particular principal diagnosis be identified Another commenter questioned the appropriateness of some exclusions and suggested postponement of implementation of the proposal until additional review could be completed.

Response: The creation of the CC exclusions list was a major project involving hundreds of thousands of codes. The proposed revisions were intended to be only a first step toward refinement of the CC list and should not be perceived as exhaustive. We intend to review the remaining CC and identify additional exclusions as appropriate.

That the CC exclusions list is not exhaustive should not be a barrier to its immediate implementation. In general, the criteria used for eliminating certain diagnoses from consideration as CC were intended to identify only the most obvious diagnoses that should not be considered complications of another diagnosis. In addition, having conducted further review of the CC exclusion list on the basis of comments, we believe the exclusions comport with the principles or criteria used to develop

them. Recognizing that some of the exclusions do represent judgments that might vary from one clinician to another, similar to other aspects of DRG classification issues, we welcome specific suggestions regarding future modification of the CC exclusions list. Comments may be submitted in writing through the Grouper Changes address published elsewhere in this document.

Comment: One commenter noted that many of the refinements to the CC proposed through the exclusion listing were targeted at coding problems such as coding symptoms, signs, and ill-defined conditions. The commenter suggested that such refinements be implemented through the MCE or the PROs. He also suggested that the ICD-9-CM Coordination and Maintenance Committee look into the coding issues raised in the CC exclusions.

Response: The purpose of the MCE is to verify certain information furnished on the claim-for example, open versus closed biopsy, invalid diagnosis or procedure code. Regardless of whether for example, both symptoms and a diagnosis of the same condition should be coded, their simultaneous appearance on a claim is nevertheless likely to be fully supported by evidence in the medical record. No correction to the data submitted by a hospital is necessary in that instance, and without a change to the CC listing the symptom would be treated by the Grouper as a CC in assigning the case to a DRG. By revising the CC list as proposed, the Grouper would no longer treat the symptom as a CC. In addition, we do not believe our FIs are necessarily the appropriate bodies to conduct coding instruction for hospitals. Yet attempting to incorporate coding rules into the MCE would require that they do so. Moreover, the issue here is not whether the coding of a claim is accurate or not. The issue is the DRG to which each claim, however coded, is assigned. And DRG assignment is performed by Grouper software, not by the MCE.

We agree that many of the refinements to the CC listing proposed were intended, to a limited extent, to serve as coding edits. That is, if a hospital reported a principal diagnosis of acute epiglottitis without obstruction. then acute epiglottitis with obstruction would not qualify as a CC for payment purposes. Many other exclusions, however, are not directly related to coding rules, such as the reporting of both acute/chronic conditions. In some cases, it is correct coding to report both the acute and chronic conditions; however, the chronic condition would not be considered a CC during an acute

manifestation. As the commenter points out, exclusions intended to represent coding edits could be implemented through the MCE or the PRO. However, we still would need to modify the Grouper program to handle the other exclusions.

Furthermore, use of the MCE could result in significant claims processing delays, as many claims would be returned to hospitals for corrections. We believe it is preferable to implement both types of exclusions through a single mechanism. Consequently, we have not adopted the commenter's suggestion.

With regard to the role of the ICD-9-CM Coordination and Maintenance Committee, the Committee is responsible for revising the ICD-9-CM and clarifying coding guidelines. We have informally asked the Committee staff to review the exclusion list and advise us of any difficulties noted. We intend to involve the Committee to the extent possible in ongoing maintenance and revision of the listing.

Comment: Three commenters noted an error in the discussion of the principles used in developing the CC refinement. These commenters advised us that coding guidelines permit the coding of both acute and chronic manifestations of the same condition but require that the acute manifestation be reported as the principal diagnosis.

Response: We regret the misstatement of coding guidelines in the proposed notice. The principles stated were intended to explain our rationale for excluding certain diagnoses as CC from other specific principal diagnoses, rather than to delineate coding principles. The commenters are correct in stating that coding guidelines permit the coding of both acute and chronic manifestations of the same condition.

Item 1 of Section V.B on page 18886 of the May 19, 1987 Federal Register should read:

 Chronic and acute manifestation of the same condition should not be considered as CC for one another.

We should also point out that correct coding guidelines require that in cases where both chronic and acute manifestations of the same disease appear and lead to the admission, the acute condition is to be reported as the principal diagnosis. PROs are responsible for verifying that Medicare bills are correctly coded based on documentation in the medical record. The CC listing does not serve this function.

#### D. Surgical Hierarchies

In the May 19, 1987 notice, we advised that we were proposing to change

surgical hierarchies. However, since those proposed changes were based on recalibration of the DRGs, they were discussed in the June 10, 1987 proposed rule, at 52 FR 220804.

We proposed to reorder the surgical hierarchies in the following MDCs as described below:

 In MDC 2, we proposed to order Orbital Procedures before Retinal Procedures.

We received no specific comments on this proposal and, therefore, are adopting it without change.

2. In MDC 3, we proposed to reorder the procedure groups as follows: Major Head and Neck Procedures Tonsil and Adenoid Procedures Except Tonsillectomy and/or Adenoidectomy Only

Cleft Lip and Palate Repair Sialoadenectomy Myringotomy with Tube Insertion Sinus and Mastoid Procedures Salivary Gland Procedures Except Sialoadenectomy

Miscellaneous Ear, Nose and Throat Procedures Rhinoplasty

Tonsillectomy and/or Adenoidectomy
Only

Other Ear, Nose and Throat O.R. Procedures

Comment: We received one comment questioning the appropriateness of reordering the tonsil and adenoid procedures except tonsillectomy and/or adenoidectomy only to the second position and reordering myringotomy with tube insertion above sinus and masterial procedures.

mastoid procedures. Response: We regrouped discharges from the FY 1986 MEDPAR file used for recalibration with the proposed DRG changes, including surgical hierarchy revisions, reflected in the Grouper. We continue to find tonsil and adenoid procedures except tonsillectomy and/or adenoidectomy only (DRGs 57 and 58) to be more resource intensive than all other surgical groups in MDC 3 except major head and neck procedures (DRG 49) and other ear, nose and throat OR procedures (DRG 63). As we explained in the proposed rule, the "other procedures" group in each MDC that has such a group consists of the procedure least directly related to the diagnoses in that MDC and is, accordingly, always ordered last in the surgical hierarchy. We believe that DRGs 57 and 58 are as resource intensive as they are because many cases assigned to them entail multiple procedures, such as tonsillectomy or adenoidectomy plus any other OR procedure. Therefore, we are ordering tonsil and adenoid

procedures except tonsillectomy and/or adenoidectomy only below major head and neck procedures as proposed.

The commenter's observation regarding the relative positions of myringotomy (DRGs 61 and 62) and sinus and mastoid procedures (DRGs 53 and 54) are borne out by analysis of data regrouped with Grouper software incorporating the change in the hierarchy.

As one can see from Table 5 of the June 10 proposal, the weighting factor for DRG 61 is greater than that for DRG 53. In point of fact, we weighted by frequency of cases in DRGs 61 and 62 for comparison to DRGs 53 and 54, as described in the proposed rule, but since DRGs 54 and 62 are low-volume DRGs, their relative weights had no practical effect on the comparison. It was on the basis of this comparison that we proposed to reorder myringotomy and sinus and mastoid procedures in the surgical hierarchy. Similar comparisons informed our other proposals regarding the surgical hierarchies.

When myringotomy preceded sinus and mastoid procedures in the hierarchy, however, the weighted average of the relative weights for DRGs 53 and 54 exceeded that for DRGs 61 and 62. On the basis of these findings, we are revising the ordering of sinus and mastoid procedures and myringotomy with tube insertion. The MDC 3 surgical hierarchy is therefore revised as follows:

Major Head and Neck Procedures (DRG 49)

Tonsil and Adenoid Procedures Except Tonsillectomy and/or Adenoidectomy only (DRGs 57-58)

Cleft Lip and Palate Repair (DRG 52) Sialoadenectomy (DRG 50) Sinus and Mastiod Procedures (DRGs

Myringotomy with Tube Insertion (DRGs 61-62)

Salivary Gland Procedures Except Sialoadenectomy (DRG 51) Miscellaneous Ear, Nose and Throat

Miscellaneous Ear, Nose and Throa Procedure (DRG 55)

Rhinoplasty (DRG 56)

53-54)

Tonsillectomy and/or Adenoidectomy Only (DRGs 59-60)

Other Ear, Nose and Throat OR Procedures (DRG 63)

3. In MDC 5, we proposed to reorder the procedure groups as follows:

Heart Transplant

Cardiac Valve Procedure with Pump Coronary Bypass

Other Cardiothoracic Procedures Major Reconstructive Vascular

Procedures
Permanent Cardiac Pacemaker
Implantation

Amputation Except Upper Limb and Toe

Vascular Procedures Except Major Reconstructive Procedures

Amputation Upper Limb and Toe Cardiac Pacemaker Replacement and/or Revision

Vein Ligation and Stripping Other Circulatory System OR Procedures

We received no specific comment on this proposal, so we are implementing it without change.

4. In MDC 6, we proposed to reorder the procedure groups as follows:

Stomach, Esophageal and Duodenal Procedures

Rectal Resection

Major Small and Large Bowel Procedures

Procedures
Peritoneal Adhesiolysis
Appendectomy

Minor Small and Large Bowel Procedures

Mouth Procedures
Anal and Stomal Procedures
Hernia Procedures

Other Digestive System OR Procedures

Comment: One commenter expressed concern about ordering stomach, esophageal and duodenal procedures above major small and large bowel procedures.

Response: As with all other hierarchy changes, we based our proposal on a comparison of the average relative weight for cases involving stomach, esophageal and duodenal procedures (DRGs 146-147) and major small and large bowel procedures (DRGs 148-149). and so on through all the procedure groups in MDC 6. While the relative weights in Table 5 of the final rule of changes to the inpatient hospital prospective payment system and FY 1988 rates have changed somewhat from those in the proposed notice, owing to use of more complete FY 1986 data, the proposed ordering of the surgical hierarchy is entirely consistent with the relative weights for the DRGs in MDC 6. Accordingly, we are implementing the proposed hierarchy changes in MDC 6 without change.

5. In MDC 8, we proposed to reorder the procedure groups as follows:

Bilateral or Multiple Major Joint Procedures of the Lower Extremity Wound Debridement and Skin Graft Except Hand

Major Joint and Limb Reattachment
Procedures

Hip and Femur Procedures Except Major Joint

Amputations
Back and Neck Procedures
Biopsies

Lower Extremity and Humerus
Procedures Except Hip, Foot, Femur

Major Shoulder/Elbow Procedures or Other Upper Extremity Procedures with CC

Knee Procedures Soft Tissue Procedures

Arthroscopy

Local Excision and Removal of Internal Fixation Devices Except Hip and Femur

Local Excision and Removal of Internal Fixation Devices of Hip and Femur

Major Thumb or Joint Procedures or Other Hand or Wrist Procedures with CC

Foot Procedures

Shoulder, Elbow or Forearm Procedures Except Major Joint Procedures without CC

Hand or Wrist Procedures Except Major Joint Procedures without CC Other Musculoskeletal System and

Connective Tissue OR Procedures We received no specific comments on this proposal. As described in the June 10, 1987 proposed rule and the foregoing discussion, however, when we propose changes to the surgical hierarchy, we are not always able to test the effects of the revisions due to the unavailability of revised Grouper software at the time of publication. Rather, in performing analysis of the surgical hierarchies, we simulate most major classification changes to approximate the placement of cases under the proposed reclassification and then recalibrate the DRG weights. The weighting factor for each procedure group then serves as our best estimate of relative resource use for

that procedure group. As occurred last year for MDC 7, when we received a revised Grouper program and were able to test the proposed hierarchy changes, we found that the revision to the surgical hierarchy in MDC 8 produced anomalous results. The proposed hierarchy changes are consistent with the DRG weights from the top of the hierarchy through knee procedures (DRGs 221-222) and from foot procedures (DRG 225) through the bottom of the hierarchy. However, there appears to be a substantial number of cases involving surgical procedures from more than one of the groups for soft tissue procedures (DRGs 226-227) arthroscopy (DRG 230), local excision and removal of internal fixation devices (DRGs 230-231), and major thumb or joint procedures (DRG 228). Similar to the diagnostic procedure group in MDC 7, we found that the number of patients with arthroscopy more than doubled when the procedure group was moved higher in the surgical hierarchy. This result indicates that arthroscopy is as frequently performed in conjunction

with a procedure from one of the abovementioned categories as it is by itself.

The fact that DRG 232 picked up so many cases, in and of itself is not troubling. However, the reassignment of so many cases results in weighting factors that are no longer appropriate for the proposed surgical hierarchy. For example, when arthroscopy is ordered 17th, as in the current DRG classification, its weight is the 12th highest in MDC 8. When arthroscopy is ordered 12th, as proposed, its weight drops to the 15th lowest. Most of the case movement seems to have come from DRGs 230 and 231 (Removal of Internal Fixation Devices, Hip and Femur, and Except Hip and Femur, respectively). Similarly to what occurred in MDC 7 last year when we proposed surgical hierarchy changes, we are left with the anomalous situation in which we cannot achieve complete correspondence between the ordering of the procedure groups in the surgical hierarchy and the relative weights that result from classification of cases based on such ordering.

Consequently, we have decided not to revise the ordering for that section of the MDC 8 surgical hierarchy involving removal of internal fixation devices, soft tissue, major thumb, and arthroscopy procedures, rather than to proceed with changes proposed for this section. We will, however, implement the other proposed changes in the MDC 8 hierarchy. Thus, the surgical hierarchy for MDC 8 is modified as follows:

Bilateral of Multiple Major Joint Procedures of the Lower Extremity (DRG 471)

Wound Debridement and Skin Graft Except Hand (DRG 217)

Major Joint and Limb Reattachment Procedures (DRG 209)

Hip and Femur Procedures Except Major Joint (DRGs 210–212) Amputations (DRG 213)

Back and Neck Procedures (DRGs 214-215)

Biopsies (DRG 216)

Lower Extremity and Humerus Procedures Except Hip, Foot, Femur (DRGs 218-220)

Major Shoulder/Élbow Procedures or Other Upper Extremity Procedures with CC (DRG 223)

Knee Procedures (DRGs 221–222)
Local Excision and Removal of Internal
Fixation Devices of Hip and Femur

(DRG 230)
Local Excision and Removal of Interal
Fixation Devices Except Hip and
Femur (DRG 231)

Soft Tissue Procedures (DRGs 226-227)
Major Thumb or Joint Procedures or
Other Hand or Wrist Procedures with
CC (DRG 228)

Arthroscopy (DRG 232) Foot Procedures (DRG 225)

Shoulder, Elbow or Forearm Procedures Except Major Joint without CC (DRG 224)

Hand or Wrist Procedures Except Major Joint Procedures without CC (DRG 229)

Other Musculoskeletal System and Connective Tissue OR Procedures (DRGs 233-234)

6. In MDC 11, we proposed to order Minor Bladder Procedures above Prostatectomy.

Comment: One commenter questioned the appropriateness of ordering minor bladder procedures (DRGs 308–309) above prostatectomy (DRGs 306–307), arguing that major procedures should be placed above, not below, minor ones.

Response: With respect to the commenter's argument that major procedures should always be placed above minor ones, we agree insofar as we are making appropriate comparisons. Were minor bladder procedures to be ordered above major bladder procedures, for example, it would call into question our use of the terms major and minor and would suggest we reconsider the relative complexity of the two groups.

On the other hand, we also believe that it is not generally appropriate simply to compare the adjectives "major" and "minor" without regard to the subjects they modify. In the instant case, "major" modifies procedures on one organ (the prostate) and "minor" is used to describe a set of procedures on another organ (the bladder). Unless we can assume that the bladder and the prostate are entirely comparable with respect to ease of surgical access. effects of surgical intervention on anatomically proximal organs, incidence of post-operative complications, and other factors, there is no basis for comparing minor procedures on the bladder to major procedures on the prostate and arguing that the latter should always precede the former. It is also important to recognize that the bladder procedures included in DRGs 308 and 309 are minor only in comparison to the bladder procedures included in DRGs 303-305.

Indeed, based on further analysis and more complete MEDPAR data than was available at the time we prepared the proposed rule, we continue to find that the weighted average of the relative weights for minor bladder procedures (DRGs 308–309) exceeds that for prostatectomy (DRGs 306–307). Therefore, we are implementing our proposed revision to the surgical hierarchy of MDC 11 without change.

In addition to the comments on specific surgical hierarchy changes proposed, discussed above, we received a comment from ProPAC reiterating its belief that the surgical procedures within each group should be evaluated on a regular basis. ProPAC believes that we did not address that part of its recommendation and that we misrepresented its position regarding the role of clinical input. ProPAC does not maintain that clinicians should determine the ordering of the surgical groups in the hierarchies, but rather recommends that clinical input should be combined with empirical analysis to produce revised procedure groups reflective of current technology and

Response: We agree with ProPAC that clinical input should be combined with empirical analysis in any broad-based revision of the procedure groups. However, we are not persuaded that there is a need for such broad-based revision of the procedure groups at the present time. We are certainly willing to conduct appropriate analysis regarding membership of the procedure groups when a classification problem comes to our attention. In the absence of complaint or concern regarding the classification of particular kinds of cases, however, we are not convinced that our limited staff resources are best utilized on a project that is tantamount to reinventing the surgical DRGs. In addition, since the procedures assigned to many procedure groups are determined by reference to specific organs within an MDC, it is not clear what alternatives ProPAC believes ought to be considered or what classification problems major evaluative work should be designed to solve.

#### V. Proposed Changes to Reduce Inappropriate DRG 468 Assignment

#### A. Background

DRG 468 (Unrelated OR procedures) is reserved specifically for those cases in which none of the surgical procedures furnished to a patient is related to the principal diagnosis. It was established as a means of identifying those cases that do not readily lend themselves to classifications within groups of clinically similar patients, because the cases themselves do not reflect typical treatment patterns. These include, for example, cases in which the patient develops pressing medical-surgical needs related to a secondary diagnosis or complication. DRG 468 is not a catchall for cases that do not fit elsewhere. It is designed to preserve the utility of,

rather than violate the principle of, diagnosis related classifications.

#### B. Reassignment of Intra-abdominal Hemangioma to MDC 6

In the May 19 notice, we proposed assigning diagnosis code 228.04, Intraabdominal hemangioma, to MDC 6, Diseases and Disorders of the Digestive System. We also proposed assigning cases treated surgically to the appropriate surgical DRG based on the site of the lesion. Cases treated medically would be assigned to DRGs 182, 183, and 184 (Esophagitis, Gastroenteritis, and Miscellaneous Digestive Disorders). We did not receive any comments on this proposal, so it is adopted without change.

#### C. Removal of Codes for Minor Skin Procedures From Surgical List

We proposed to remove procedure codes 86.09, Other incision of skin and subcutaneous tissue, and 86.3, Other local excision or destruction of lesion or tissue of skin and subcutaneous tissue, from the list of OR procedures. We did not receive any comments on this proposal, so it is adopted without change.

#### D. Adding Lymphatic Structure Biopsy to MDC 1

We proposed adding procedure code 40.11, Biopsy of lymphatic structure, to MDC 1, DRGs 7 and 8, Peripheral and Cranial Nerve and Other Nervous System Procedures with CC and without CC, respectively. We did not receive any comments on this proposal, so it is adopted without change.

#### E. Adding Total Splenectomy to MDC 5

We proposed adding procedure code 41.5, Total splenectomy, to MDC 5, DRG 120 (Other Circulatory System OR Procedures). We did not receive any comments on this proposal, so it is adopted without change.

#### F. Adding Certain Pancreas Procedure Codes to MDC 10

We proposed to add the following procedure codes to MDC 10, DRGs 292 and 293 (Other Endocrine, Nutritional and Metabolic Procedures).

- 52.2 Local excision or destruction of pancreatic lesion
- 52.51 Proximal pancreatectomy
- 52.52 Distal pancreatectomy
- 52.53 Radical subtotal pancreatectomy
- 52.59 Other partial pancreatectomy, not elsewhere classified

We did not receive any comments on this proposal, so it is adopted without change.

#### G. MDC 11 Issues

We proposed adding procedure code 70.77, Vaginal suspension and fixation, to MDC 11, DRGs 308 and 309, (Minor Bladder Procedures). We also proposed adding procedure codes for replacement and removal of penile prosthesis, 64.95 through 64.97 to MDC 11, DRG 315 (Other Kidney and Urinary Tract OR Procedures). Finally, we proposed adding procedure code 60.69, Other Prostatectomy to DRGs 306 and 307 (Prostatectomy). We did not receive any comments on these proposals, so they are adopted without change.

# H. Adding a Urethral Repair Code to MDC 12

We proposed adding procedure code 58.49, Other Urethral Repair, not elsewhere classified, to MDC 12, DRG 341, (Penis Procedures) to prevent inappropriate assignment of cases to DRG 468 when the procedure is related to the principal diagnosis. We did not receive any comments on this proposal, so it is adopted without change.

#### I. Deletion of Certain Codes from the OR List

We proposed deletion of procedure codes, 39.62, Hypothermia systemic incidental to open heart surgery, 39.63, Cardioplegia, and 39.64 Intraoperative cardiac pacemaker, from the list of operating room procedures. We did not receive any comments on this proposal, so it is adopted without change.

# VI. Comments and Responses on Coding Issues

#### A. Background

In the final notice on changes to the DRG classification system published June 3, 1986 (51 FR 20192), and the final rule on the prospective payment system published September 3, 1986 (51 FR 31454), we published lists of new ICD-9-CM codes that became effective October 1, 1986. These new codes were adopted as a result of the recommendation of the ICD-9-CM Coordination and Maintenance Committee, and were also published on August 29, 1986 (51 FR 30914) in a notice announcing all codes approved before July 1, 1986, the availability of related instructional material, additions to the ICD-9-CM indexes, and errata for volumes 1, 2, and 3 of ICD-9-CM. Both our May 19 notice and this final notice contain all new coding changes that were approved before July 1, 1987 since the ICD-9-CM Coordination and Maintenance Committee does not plan to publish a separate listing this year.

Comment: Two commenters wrote expressing concern with the process for

addressing coding issues. The commenters specifically encouraged increased involvement of coding experts in the process and acceleration of the time associated with implementing new codes.

Response: We have been quite pleased with the improvements in ICD-9-CM coding that have taken place over the past two years. We feel the committee has taken a giant step in improving coding in the hospital industry.

Although formal membership on the committee has not been extended to non-government entities, we note that coding experts from the American Hospital Association, the American Medical Records Association, the Commission on Professional and Hospital Activities and numerous other organizations have attended every meeting of the committee and have provided extensive recommendations and comments on proposed revisions. The insights, experiences and ideas shared by these participants have contributed significantly to the coding developments and have influenced the committee's recommendations. The Department of Health and Human Services is currently evaluating a proposal to add industry representatives to the committee; and a decision is expected in the near future.

Despite the fact that these commenters have recommended acceleration of the process for coding improvements, several other members of the public have expressed concern both formally and informally with the rapid pace of coding changes. Many users find it extremely difficult to make the rapid changes necessary to software programs that are used for numerous purposes to accommodate the implementation of new codes. Given the operational problems that rapid coding changes pose for hospitals, third party payers, and others, we do not believe it is appropriate to make coding changes any more frequently than annually.

Finally, we are not convinced that there is a large backlog of coding issues awaiting the committee's attention. Reactions from the American Medical Record Association and others indicate that the committee has made dramatic steps in resolving longstanding coding problems. However, the committee is evaluating means of improving its revision process, including the possibility of revising one or more chapters. The public is invited to raise issues that have not been included on the agenda at each meeting. If commenters have coding items that they believe require attention of the

committee, they are encouraged to submit a detailed request to the committee co-chairperson whose address is identified elsewhere in this document, identifying the problem, pertinent background information and recommended solutions.

#### B. Proposed Removal of Certain Codes from the Surgical List

We proposed to remove the following procedure codes from the list of surgical procedures:

38.22 Percutaneous angioscopy 44.22 Endoscopic dilation of pylorus 44.93 Insertion of gastric bubble

(balloon)

44.94 Removal of gastric bubble (balloon)

51.97 Therapeutic endoscopic
 procedures on biliary tract, oral route
 51.98 Other percutaneous procedures
 on biliary tract

55.03 Percutaneous nephrostomy without fragmentation

55.04 Percutaneous nephrostomy with fragmentation

80.52 Intervertebral chemonucleolysis Thus, the presence of any one of these procedure codes would not result in assignment of a case to a surgical DRG.

Comment: Three commenters from a single hospital wrote to express displeasure with the removal of the chemonucleolysis procedure, 80.52, from the list of surgical procedures. They maintain that the necessity of sterile conditions and the risk of anaphylactic shock make it unsafe to perform the procedure in settings other than an OR.

Response: We recognize that an operating room may be the setting of choice for some facilities that perform chemonucleolysis. However, our medical consultants have advised us that the procedure is non-invasive and may safely be performed in settings other than ORs with proper precautions.

We note that nearly any procedure involving an injection presents the possibility of anaphylactic shock. It can be reasonable for a hospital to choose to use an operating room for prophylactic reasons in situations presenting even a small risk of anaphylactic shock. According to the commenters, one percent of patients develop such shock. However, we do not believe this is justification to place a procedure on the OR list. We note that this list is intended to recognize surgical procedures rather than procedures that are carried out in an operating room for prophylactic reasons.

Finally, if we were to place procedures on the OR list because of the possibility of anaphylactic shock, nearly all radiographic procedures and many other non-surgical procedures would need to be reclassified to the OR list. We should point out that the DRG classification of a procedure is not intended to influence the practice patterns at any particular hospital. Hospitals should continue to choose appropriate settings for procedures based upon the conditions at the facility and the individual needs of their patients.

Comment: Numerous commenters wrote expressing concern with our proposal to remove procedure codes 55.03 and 55.04, Percutaneous nephrostomy, without and with fragmentation, respectively, from the list of OR procedures. These commenters noted the necessity of general anesthesia and sterile conditions, protracted procedure time, as well as the possibilities of severe complications and profuse blood loss as contraindications for using other settings, such as radiographic suites. All commenters urged that we reconsider our proposal. One of the commenters noted that the current classification structure may result in payments higher than cost in percutaneous lithotripsy cases involving simple stone removal, but that complicated stone removal which allegedly comprises the majority of cases, was underpaid. Consequently, an alternative classification structure was recommended.

Response: Based on the number and nature of the comments received in response to this proposal, we examined preliminary FY 1987 data on cases involving percutaneous nephrostomy. We are persuaded that the data on resource intensity (in terms of both standardized charges and length of stay) support the commenters' arguments that moving cases involving percutaneous nephrostomy from the surgical to the medical DRGs would result in systematic underpayment of such cases. Accordingly, we believe it is appropriate to defer action at this time. Percutaneous nephrostomy with and without fragmentation will remain on the list of operating room procedures as currently assigned and will result in classification of cases to DRGs 303, 304, 305, 442 and 443. We will further evaluate this classification issue in light of alternatives suggested by commenters as more data on the procedure become available. Any further proposal will be published for additional public comment.

# C. Proposed Removal of a Code From the CC List

We proposed removing diagnosis code 795.8, Positive serological or viral culture finding for Human T-Cell Lymphotropic Virus-III/ Lymphadenopathy-Associated Virus (HTLV-III/LAV) from the list of CC. We received no negative comments and ProPAC's support for this proposal, so it is adopted without change.

#### D. New Coding Changes

We notified the public of plans to add new and revised diagnosis and procedure codes and the deletion of several procedure codes. Since the ICD-9-CM Coordination and Maintenance Committee does not plan to publish a separate listing this year, we are republishing the tables in this final notice. Table 1 contains the new or revised ICD-9-CM diagnosis codes, Table II lists the new or revised ICD-9-CM procedure codes, Table III lists the revised pacemaker DRG logic tables, and Table IV lists the ICD-9-CM procedure codes that we are deleting. A copy of the ICD-9-CM Official Authorized Addendum will be provided to each hospital by the FI in September 1987. Additional copies will be available from the Government Printing Office, in the September issue of the Journal of American Medical Record Association, and in Coding Clinic for ICD-9-CM.

Comment: One commenter expressed concern with the proposed DRG assignment of procedure code 86.06. Insertion of infusion pump. The commenter believes the proposed classification does not adequately recognize use of the pump in treatment of patients with osteomyelitis or in delivery of morphine for terminally ill patients. The commenter suggested assignment to DRGs 4 and 210.

Response: Procedure code 86.06 has been assigned to the other operating room procedures DRG of nearly every MDC that contains such a group. Only MDCs 2, 3 and 5 have been omitted, due to the unlikelihood of the device being used in treatment of patients with eve. ear or circulatory system diagnoses. Patients with a principal diagnosis of osteomyelitis would be assigned to DRGs 233 or 234, based on presence or absence of CC, if implantation of the infusion pump were the only surgical procedure performed. If another MDC 8 surgical procedure were performed in addition to insertion of an infusion pump, the surgical hierarchy would base DRG assignment on the other surgical procedure.

Similarly, patients being treated for pain associated with terminal illness would be assigned to the other OR procedure DRG for the MDC in which the principal diagnosis falls, if implantation of the infusion pump is the only surgical procedure performed.

Since the other OR procedure group is uniformly considered last in the surgical hierarchy, DRG assignment of cases involving multiple procedures within the MDC will be based on any other surgical procedure performed that is

related to the principal diagnosis rather than on the implantation of an infusion

We find no problem with the proposed classification of implantation of the infusion pump. It appears the

commenter's concern is based on an oversight in reading the list of DRGs to which the procedure is assigned or a misunderstanding of the classification mechanism.

#### TABLE I.—NEW OR REVISED DIAGNOSIS CODES

Diagnosis code 1	Description	MDC	DRG ²
996.53 996.54 996.59 996.80 996.81 996.82 996.83 996.84	Respiratory failure Other pulmonary insufficiency, not elsewhere classified. Other diseases of lung, not elsewhere classified and an elsewhere classified and an elsewhere classified and an elsewhere classified and an elsewhere classified and elsewhere classified.  Mechanical complication due to graft of other tissue, not elsewhere classified.  Mechanical complication due to ocular lens prosthesis.  Mechanical complication due to breast prosthesis.  Mechanical complication due to other implant and internal device, not elsewhere classified.  Complications of transplanted organ, not otherwise specified.  Complications of transplanted kidney Complications of transplanted liver.  Complications of transplanted heart.  Complications of transplanted lung Complications of transplanted pancreas	21 9 91	87. 99 and 100. 101 and 102. 101 and 102. 46, 47, and 48. 452 and 453. 46, 47, and 48. 276. 452 and 453. 452 and 453. 331, 332, and 333 205 and 206. 144 and 145. 101 and 102.
996.86 996.89	Complications of transplanted particless	21	204. 452 and 453.

¹ All of the new diagnosis codes, except 518.89, would be added to the list of CC.

² DRG listed is assignment based on non-surgical treatment. If an OR procedure is performed, DRG assignment within the MDC is determined by the OR procedure performed.

3 Not added to the list of CC.

#### TABLE II.—New OR REVISED PROCEDURE CODES

Proce- dure code	Description	DRG
01.11	Closed [percutaneous] [needle] biopsy of cerebral meninges	non-OR.
01.12	Open biopsy of cerebral meninges	1, 2, 3, 400, 406, and 407.
01.13	Closed [percutaneous] [needle] biopsy of brain.	
01.14	Open biopsy of brain	1, 2, 3, 400, 406, and 407.
03.90	Insertion of catheter into spinal canal for infusion of therapeutic or palliative substances	
04.11	Closed [percutaneous] [needle] biopsy of cranial or peripheral nerve or ganglion	non-OR.
04.12	Open biopsy of cranial or peripheral nerve or ganglion	
	open sope, or called or perpendicular or gangiori	442, and 443.
06.11	Closed [percutaneous] [needle] biopsy of thyroid gland	
06.12	Open biopsy of thyroid gland	290.
07.11	Closed [percutaneous] [needle] biopsy of adrenal gland	non-OR.
07.12	Open biopsy of adrenal gland	286.
27.22	Biopsy of uvula and soft palate	63, 168, and 169
33.24	Closed biopsy [endoscopic] of bronchus	non-OR.
33.25	Open biopsy of bronchus	75.
33.26	Closed percutaneous [needle] biopsy of lung	non-OR.
33.27	Closed endoscopic biopsy of lung	76 and 77.
33.28	Open biopsy of lung	75, 233, 234, 315
		400, 406, and 407.
33.29	Other diagnostic procedures on lung and bronchus	
34.25	Closed [percutaneous] [needle] biopsy of mediastinum	non-OR.
34.26	Open biopsy of mediastinum	76, 77, 292, 293,
		394, 400, 406,
36,01	Single vessel percutaneous transluminal coronary angioplasty [PTCA] without mention of thrombolytic agent	
36.02	Single vessel percutaneous transluminal coronary angioplasty [PTCA] with thrombolytic agent	108 and 112.
36.05	Multiple vessel percutaneous transluminal coronary angioplasty [PCTA] performed during the same operation with or without mention of thrombolytic agent.	108 and 112.
37.70	Initial insertion of lead [electrode], not otherwise specified.	(1)

# TABLE II.—NEW OR REVISED PROCEDURE CODES—Continued

Proce- dure code	Description	DRG
37.71	Initial insertion of transvenous lead [electrode] into ventricle	(1)
37.72	Initial insertion of transvenous leads [electrodes] into atrium and ventricle	(ii)
37.73	Initial insertion of transversors lead felectrods] into attain and vertice	. (7)
	Initial insertion of transvenous lead [electrode] into atrium	. (1)
37.74	Insertion or replacement of epicardial lead [electrode] into epicardium	. (1)
37.75	Revision of lead [electrode]	. 117, 442, and
		443.1
37.76	Replacement of transvenous atrial and/or ventricular lead(s) [electrode]	. (1)
37.77	Removal of lead(s) [electrode] without replacement	117 442 and
		443.1
37.78	Inserting of temporary transposes assembles existen	
	Insertion of temporary transvenous pacemaker system	
37.79	Revision or relocation of pacemaker pocket	
		443.1
37.80	Insertion of permanent pacemaker, initial or replacement, type of device not specified	. (1)
37.81	Initial insertion of single-chamber device, not specified as rate responsive	. (1)
37.82	Initial insertion of a single chamber device, rate responsive	(1)
37.83	Initial insertion of dual chamber device	100
37.85	Replacement of any type pacemaker device with single-chamber device, not specified as rate responsive	
37.86	Poplacement of any type pacernane device with single-chamber device, not specified as rate responsive	
	Replacement of any type pacemaker device with single-chamber device, rate responsive	. (*)
37.87	Replacement of any type pacemaker device with dual-chamber device	. (1)
37.89	Revision or removal of pacemaker device	. 117, 442, and
		443.1
41.32	Closed [aspiration] [percutaneous] biopsy of spleen	non-OR.
41.33	Open biopsy of spleen	392, 393, 400.
47.00	Open propagation	
44.14	Clearly forders and the second	406, 407.
	Closed [endoscopic] biopsy of stomach	. non-OR.
44.15	Open biopsy of stomach	. 154, 155, and 156.
45.14	Closed [endoscopic] biopsy of small intestine	non-OR.
45.15	Open biopsy of small intestine	152 and 153.
45.25	Closed [endoscopic] biopsy of large intestine.	non-OR
45.26	Ones higher of large intecting	150 and 150
	Open biopsy of large intestine	. 152 and 153.
45.95	Anastomosis to anus	. 148, 149, 400,
27000		406, 407, 442,
		and 443.
48.24	Closed [endoscopic] biopsy of rectum	non-OR.
48.25	Open biopsy of rectum	152 and 153
50.11	Closed (percutaneous) [needle] biopsy of liver	non-OR.
50.12	Open biopsy of liver	60 76 77 170
00.12	Open biopsy of five	
111111111111111111111111111111111111111		171, 199, 200,
98		233, 234, 269,
-		270, 292, 293,
- 120		315, 344, 345,
133900		365, 394, 400,
- 17.00	Participation of the second of	406, 407, 442,
3 0 00		and 443.
51.12	Closed [percutanguis] biopsy of callbladder as hills dusts	
51.13	Closed [percutaneous] biopsy of gallbladder or bile ducts	170 171 100
31.13	Open biopsy of gallbladder or bile ducts	
20		200.
52.11	Closed [aspiration] [needle] [percutaneous] biopsy of pancreas	non-OR.
52.12	Open biopsy of pancreas	. 170, 171, 199,
		200, 292, 293,
J 1912 E.		400, 406, 407,
- Cla		
64.04	Closed Engravianous Foodle News of the state	442, and 443.
54.24	Closed [percutaneous] [needle] biopsy of intra-abdominal mass	. non-OR.
55.23	Closed [percutaneous] [needle] biopsy of kidney	. non-OR.
55.24	Open biopsy of kidney	. 233, 234, 303,
1		304, 305, 394,
- You have		400, 406, 407,
		442, and 443.
56.32	Closed percutangous highest of ureter	
120000000	Closed percutaneous biopsy of ureter	non-OR.
56.33	Closed endoscopic biopsy of ureter	. non-OR.
56.34	Open biopsy of ureter	. 303, 304, and 305.
56.35	Endoscopy [cystoscopy] [looposcopy] of ileal conduit	. non-OR.
57.33	Closed [transurethral] biopsy of bladder	310 311 344
CHAIR STREET		
57.24	Open bioppy of bladder	345, and 365.
57.34	Open biopsy of bladder	
Marie -		345, 365, 400,
		406, and 407.
-	Closed [percutaneous] [needle] biopsy of prostate	

# TABLE II.—NEW OR REVISED PROCEDURE CODES—Continued

Proce- dure code	Description	DRG
60.12	Open biopsy of prostate	
60.13	Closed [percutaneous] biopsy of seminal vesicles	345.
60.14	Open biopsy of seminal vesicles	non-OR. 344 and 345.
62.11	Closed [percutaneous] [needle] biopsy of testis	non-OR.
62.12	Open biopsy of testis	338, 339, and 340
68.13	Open biopsy of uterus	
68.14	Open biopsy of uterine ligaments	358, and 359. 354, 355, 357,
00.45		358, and 359.
68.15	Closed biopsy of uterine ligaments	
78.40	Closed biopsy of uterus	363 and 364. 233, 234, 442, and
, 0, 10	outer repair and places operations on bone, unspecified site	443.
78.41	Other repair and plastic operations of chest cage	76, 77, 233, 234,
		442, and 443.
78.42	Other repair and plastic operations of humerus	COMPANY OF THE PARTY OF THE PAR
78.43	Other reads and plastic apprehing of realism (slave	442, and 443.
10.43	Other repair and plastic operations of radius/ulna	223, 224, 442, and
78.44	Other repair and plastic operation of carpals/metacarpals	CONTRACTOR
78.45	Other repair and plastic operations of femur	
		442, and 443.
78.46	Other repair and plastic operations of patella.	221, 222, 442, and
70.47		443.
78.47	Other repair and plastic operations of tibia/fibula	
78.48	Other repair and plastic operations of tarsals/metatarsals	442, and 443. 225, 442 and 443.
78.49	Other repair and plastic operation on bone, not elsewhere classified	233, 234, 442, and
	Charles and place application of solid, not discussed discussed	443.
78.90	Insertion of bone growth stimulator, unspecified site	
The same of the sa		443.
78.91	Insertion of bone growth stimulator into chest cage	
78.92	Insertion of bone growth stimulator into humerus	442, and 443.
10.02	insertion of both grown simulator into futileus	442, and 443.
78.93	Insertion of bone growth stimulator into radius/ulna	
		443.
78.94	Insertion of bone growth stimulator into carpals/metacarpals	228, 229, and 441
78.95	Insertion of bone growth stimulator into femur	
78,96	Incortion of hone growth etimulates into notalla	442, and 443.
70.50	Insertion of bone growth stimulator into patella	443.
78.97	Insertion of bone growth stimulator into tibia/fibula	
		442, and 443.
78.98	Insertion of bone growth stimulator into tarsals/metatarsals	225, 442, and 443
78.99	Insertion of bone growth stimulator, not elsewhere classified	
85.11	Closed Ferral teneral Ferral Name of Ferral	443.
85.12	Closed [percutaneous] [needle] biopsy of breast	
05.12	Open stopey of stodet.	292, 293, 442,
	TO CALL THE STATE OF THE STATE	and 443.
85.95	Insertion of breast tissue expander.	259, 260, 261,
00.00		442, and 443.
85.96	Removal of breast tissue expander(s)	
86.93	Insertion of tissue expander	442, and 443.
00.50	modulon or ussue expander	7, 8, 63, 120, 170, 171, 217, 263,
		264, 265, 266,
		287, 439, 458,
		and 472.

TABLE II.—New OR REVISED PROCEDURE CODES—Continued

Proce- dure code	Description	DRG
86.06	Insertion of infusion pump	7, 8, 76, 77, 170, 171, 201, 233, 234, 269, 270,
		292, 293, 315, 344, 345, 365, 394, 401, 402,
99.85	Hyperthermia for treatment of cancer	408, 442, 443 459, and 472. non-OR.
99.86	Non-invasive placement of bone growth stimulator	non-OR.

¹ See pacemaker DRG logic tables following this table.

#### Table III.—Pacemaker Logic Tables

DRGs 115 and 116 currently represent and will continue to represent insertion of total pacemaker systems, that is, device plus lead(s). Although insertion of a pacemaker system used to be represented by a single procedure code (37.70, 37.73, 37.74, 37.75, 37.76 or 37.77), the revised ICD-9-CM coding system uses at least two codes to signify the insertion of a pacemaker system-one code to identify the type of device (single-chamber, dual-chamber, or rate responsive) and another to identify the type of lead(s) inserted (transvenous atrial, transvenous ventricular, or epicardial). Consequently, effective for discharges on or after October 1, 1987. the Grouper logic is modified to comport with the revised pacemaker procedure codes order for a case to be classified into DRGs 115 or 116 (Permanent Cardiac Pacemaker Implant, with AMI, Heart Failure or Shock, and without AMI, Heart Failure or Shock, respectively).

DRGs 115 or 116 will be assigned, depending on principal diagnosis, only if one of the following combinations of procedure codes appears on the claim:

37.70 and 37.80 37.70 and 37.81 37.70 and 37.82 37.70 and 37.85 37.70 and 37.86 37.70 and 37.87 37.71 and 37.80 37.71 and 37.81 37.71 and 37.82 37.71 and 37.85 37.71 and 37.86 37.71 and 37.87 37.72 and 37.80 37.72 and 37.83 37.73 and 37.80 37.73 and 37.81 37.73 and 37.82 37.73 and 37.85 37.73 and 37.86 37.73 and 37.87 37.74 and 37.80 37.74 and 37.81 37.74 and 37.82 37.74 and 37.83 37.74 and 37.85 37.74 and 37.86 37.74 and 37.87 37.76 and 37.80 37.76 and 37.80

DRG 117 is renamed Cardiac Pacemaker Revision Except Device Replacement. Procedures assigned to it include:

37.74 37.75 37.76 37.77 37.79 37.89

DRG 118 is renamed Cardiac Pacemaker Device Replacement and includes the following procedures:

37.80 37.85 37.86 37.87

Finally, the following pacemaker procedure codes will be ignored by the Grouper when they do not appear in combination with other pacemaker procedure codes as in DRGs 115 and 116 above:

37.70 37.71 37.72 37.73 37.81 37.82

These codes represent the initial insertion of a lead (37.70–37.73) or a device (37.81–37.83). We are aware of no clinical condition that would require the initial insertion of a lead without the simultaneous insertion or replacement of a device. Likewise, the initial

insertion of a device without the initial insertion of a lead(s) is clinically illogical. Accordingly, if an initial lead is inserted with no device or an initial device is inserted without a lead insertion, the procedure would not be a covered procedure since section 1862(a)(1)(A) of the Act prohibits payment for items and services that are not reasonable and necessary for the diagnosis or treatment of an illness or injury. Similarly, when an initial device is inserted but no lead code appears, a total system has not been implanted.

The following ICD-9-CM codes are deleted without replacement.

TABLE IV.—DELETED PROCEDURE CODES

Procedure code	Description
37.84	Removal of epicardial electrode:
99.71	Mercury-zinc pacemaker battery.
99.72	
99.73	
99.74	Other pacemaker battery type.
99.75	
99.76	Triggered demand pacemaker sensing type.
99.77	Inhibited demand pacemaker sensing type.
99.78	Other pacemaker sensing type.
99.79	Programmable pacemaker.

#### VII. Summary of Changes

As stated in our discussion of the comments and responses, we have made some changes to the proposals in the notice published on May 19, 1987. With the exception of the following changes, this final notice implements the proposals made in the May 19, 1987 proposed notice and the June 10 proposed rule.

## · MDC

We have decided to postpone moving diagnosis code 759.8, (Other Hamartoses, NEC) from MDC 17 to MDC 8 for FY 1988. We will study the issue in greater detail during the upcoming fiscal year and will report on our findings in the next year's proposed notice.

 Deletion of Certain Codes from the Surgical List

Based on the number and nature of the comments received, we believe it is appropriate to leave Percutaneous nephrostomy without and with fragmentation (procedure codes 55.03 and 55.04, respectively) on the list of OR procedures as currently assigned. Thus, percutaneous nephrostomy cases will continue to be assigned to DRGs 303, 304, 305, 442, and 443. We will further evaluate this classification issue in light of the alternative suggestions by commenters as more data on the procedure becomes available.

#### Surgical Hierarchy

We are reversing the proposed ordering of sinus and mastoid procedures and myringotomy with tube insertion in MDC 3.

We have decided not to revise the ordering for the section of the proposed surgical hierarchy involving removal of internal fixation devices, soft tissue, major thumb, and arthroscopy procedures.

#### Corrections

We have made some minor changes to the ICD-9-CM DRG coding changes tables and have reprinted them. Additionally, we have made minor revisions to the lists of CC refinements.

#### VIII. Regulatory Impact Statement

#### A. Executive Order 12291

Executive Order (E.O.) 12291 requires us to prepare and publish a final regulatory impact analysis for final notices such as this if the implementation of the notice meets the criteria of a "major rule". A notice is considered a major rule if its implementation is likely to result in:

(1) An annual effect on the economy

of \$100 million or more;

(2) A major increase in costs or prices for consumers, individual industries, Federal, State, or local government agencies, or geographic regions; or

(3) Significant adverse effects on competition, employment, investment, productivity, innovation, or on the ability of United States-based enterprises to compete with foreign-based enterprises in domestic or export markets.

We do not believe that any of the changes to the DRG classification system presented in this notice meet the E.O. criteria for a major rule. Accordingly, we have not prepared a final regulatory impact analysis for this notice. Instead, we refer interested readers to the regulatory impact analysis for the final rule on FY 1988 changes to the prospective payment system, which is published elsewhere in this issue of the Federal Register. In that analysis, we include the effects of all these changes to the DRG classification system in our assessment of the impact of DRG recalibration on hospitals. We also discuss there the effects of discontinuing the exclusion of alcohol and drug abuse treatment facilities from the prospective payment system.

#### B. Regulatory Flexibility Act

It is our practice to prepare and publish a final regulatory flexibility analysis that is consistent with the Regulatory Flexibility Act of 1980 (RFA) (5 U.S.C. 601 through 612) for a final notice such as this, unless the Secretary certifies that implementation of the notice will not have a significant economic impact on a substantial number of small entities. We treat all hospitals under the prospective payment system as small entities for purposes of the RFA.

As noted above, these changes to the DRG classification system will affect the amounts hospitals receive under the prospective payment system for furnishing services to Medicare beneficiaries. Therefore, this notice will clearly affect a substantial number of small entities. However, we do not consider an economic impact on small entities to be significant unless their annual total costs or revenues will be increased or decreased by at least three percent. Some of these classification changes may affect the amount paid for a particular DRG by more than 3 percent. The elimination of the age over 69 criterion, for example, may have significant effects on the weights of some DRGs as discussed elsewhere in this notice. However, the aggregate impact of these changes on hospital revenues is likely to be less substantial.

The changes we are presenting in this final notice will be used to determine the DRG weights for discharges occurring on or after October 1, 1987. However, it must be remembered that a DRG weight is a measure of average resource utilization for a particular group of cases relative to the average for all cases. Thus, each change that affects the group to which a case is assigned affects not only the payment for the

reassigned case, but the weight for all other cases in both the prior and the new group. Through annual recalibrations, all weights are readjusted to reflect all reassignments, based on the best available data. Through this process some changes are offset by others. The interaction is complex and, of course, differs from year to year. However, the end result is that most hospitals will receive higher payments for some cases and lower payments for others. Thus, we view it as highly unlikely that a substantial number of hospitals would experience increases or decreases of revenues of more than three percent solely as a result of these changes in DRG classification.

Hypothetically, a given year's classification changes could have an effect of such magnitude on some hospitals. We expect that most of these hospitals will have a high proportion of cases falling in particular strongly affected DRGs. In such instances, we believe that improvements to the classification system tend to correct systematic understatements or overstatements of average resource utilization. In the first case, those hospitals most affected would be significantly benefited. In the latter case, we would be ending an inappropriate windfall. In either case, the payment system as a whole will more closely match payments made to hospitals for Medicare-covered services with the level of resources used in providing those services.

Ultimately, we believe that these finalized changes will yield DRG groupings based on factors that better predict resource utilization than the current factors. These refinements will lead to better classification of cases within groups (where better is defined in terms of predictive power, homogeneity within groups, and differences between groups), which in turn implies better case mix measurement and improvements in case level equity (that is, payment of cases in line with their relative resource intensity) even if these changes do not significantly affect the case mix indexes of many hospitals. Thus, we see these refinements, such as eliminating the age over 69 criterion and tailoring the CC list to each principal diagnosis, as necessary first steps towards broader refinements (such as introducing severity adjustments).

For these reasons, we have determined, and the Secretary certifies.

that this final notice is not likely to have a significant economic impact on a substantial number of small entities. Therefore, we have not prepared a final regulatory flexibility analysis for this notice.

(Sections 1102, 1871, and 1886(d)(4) of the Social Security Act (42 U.S.C. 1302, 1395hh, and 1395ww(d)(4)); 42 CFR 412.10)

(Catalog of Federal Domestic Assistance Program No. 13.774, Medicare Supplementary Medical Insurance)

August 25, 1987.

William L. Roper,

Administrator, Health Care Financing Administration.

Approved: August 26, 1987.

Otis R. Bowen,

Secretary.

[FR Doc. 87-19989 Filed 8-27-87; 12:15 pm]

BILLING CODE 4120-01-M



Tuesday September 1, 1987



Part IV

# Department of Health and Human Services

Health Care Financing Administration

42 CFR Part 412

Capital Payments under the Inpatient Hospital Prospective Payment System; Final Rule

#### DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

42 CFR Part 412

[BERC-403-F]

Capital Payments Under the Inpatient Hospital Prospective Payment System

AGENCY: Health Care Financing Administration (HCFA), HHS.

ACTION: Final rule.

SUMMARY: We are amending the Medicare regulations governing the inpatient hospital prospective payment system to incorporate capital costs into that system.

EFFECTIVE DATE: This final rule is effective on October 1, 1987.

FOR FURTHER INFORMATION CONTACT: Linda Magno, (301) 594-9343.

SUPPLEMENTARY INFORMATION:

#### L. Background

In this final rule, we are changing the regulations that govern the way in which inpatient hospital capital costs, not including payments to proprietary hospitals for a return on equity capital, will be treated for Medicare payment purposes effective with hospital cost reporting periods beginning on or after October 1, 1987. Capital costs under Medicare include depreciation, interest. taxes, insurance and similar expenses (defined further in 42 CFR 413.130) for plant and fixed equipment, and for moveable equipment. Affected regulations are located in 42 CFR Part 412.

Currently, inpatient operating costs are the only costs included in the prospective payments received by hospitals under the prospective payment system (§ 412.2(c)) and by the payment amounts received under the target rateof-increase limits by hospitals and distinct part units of hospitals that are excluded from the prospective payment system (§ 413.40). Under current Medicare rules, payment for capital costs has been on a reasonable cost basis (§ 413.5) because, under section 1886(a)(4) of the Social Security Act (the Act), those costs have been specifically excluded from the definition of inpatient operating costs both for hospitals subject to the prospective payment system and for those hospitals and units excluded from that system.

With the exception of sole community hospitals, this final rule eliminates this distinction between hospital inpatient capital and inpatient operating costs for Medicare inpatient hospital services provided by hospitals subject to the prospective payment system for cost reporting periods beginning on or after October 1, 1987. Under section 1886(g)(3) of the Act, as amended by section 9303(a) of the Omnibus Budget Reconciliation Act of 1986 (Pub. L. 99–509), sole community hospitals are exempt from the inclusion of capital costs in operating costs of inpatient hospital services through their cost reporting periods beginning before October 1, 1990.

On May 19, 1987, we published a notice of proposed rulemaking (NPRM) in the Federal Register (52 FR 18840) to incorporate the capital costs of hospitals subject to the prospective payment system into that payment system. Technical corrections to the NPRM were published on June 11, 1987 (52 FR 22359). In setting forth the May 1987 NPRM, we took into consideration the provisions of section 9303 of Pub. L. 99-509, enacted on October 21, 1986, which amended section 1886(g) of the Act. As amended, section 1886(g)(3) of the Act provides that the amount of Medicare payments for capital costs attributable to inpatient services of prospective payment hospitals, otherwise determined to be reasonable, be reduced by-

 Three and one-half percent for payments attributable to portions of cost reporting periods occurring during FY 1987;

 Seven percent for payments attributable to portions of cost reporting periods or discharges (as the case may be) occurring during FY 1988; and

· Ten percent for payments attributable to portions of cost reporting periods or discharges (as the case may be) occurring during FY 1989. Section 1886(g)(3) of the Act also provides that, in any inclusion of capital costs into the prospective payment system, the aggregate Medicare capital payments, for those capital costs attributable to portions of cost reporting periods occurring during Federal fiscal years (FYs) 1988 and 1989, must be neither greater nor less than the aggregate Medicare capital payments that would have been made, taking into account the reductions mandated under section 9303 of Pub. L. 99-509, without such inclusion of capital costs into the prospective payment system.

We also considered the recommendations of the Prospective Payment Assessment Commission (ProPAC), made under the authority of section 1886(d)(4)(D) of the Act, concerning the inclusion of capital costs into the prospective payment system. Our responses to ProPAC's recommendations may be found in the May 1987 NPRM at 52 FR 18853.

In addition, section 9304 of Pub. L. 99-509 amended section 1886(d) of the Act (by adding paragraph (d)(9)) to provide for inclusion of hospitals in Puerto Rico under the prospective payment system effective with inpatient hospital discharges occurring in FY 1988. We are implementing the provisions concerning hospitals in Puerto Rico in a separate final rule. (We note that in that document we established a new Subpart K to Part 412 to implement those provisions. In the May 1987 NPRM, we proposed to add § 412.214 to Subpart K to implement the provision for incorporating capital payments into the prospective payment system for hospitals in Puerto Rico.) Section 9304 of Pub. L. 99-509 also revised section 1886(d) of the Act (by adding section 1886(d)(9)(D)(iii)) to authorize the Secretary to make exceptions and adjustments under which we believe a transition period and other refinements for incorporating capital payments into the prospective payment system could be extended to hospitals in Puerto Rico. (We note that the reductions in capital payments for FY 1988 and FY 1989, as provided under section 1886(g)(3) of the Act, also apply to hospitals in Puerto

In the May 1987 NPRM, we set forth the following proposals:

- We proposed to establish national urban and rural capital rates separately for plant/fixed equipment and for moveable equipment using the best data currently available, that is, capital costs for cost reporting periods beginning in Federal fiscal year (FY) 1984. (Since the Federal capital rates are calculated according to the Federal FY, we proposed to update the national capital rates each Federal fiscal year.)
- We proposed to standardize the capital costs for differences in case mix complexity, indirect medical education, and disproportionate share payments (and for moveable capital costs, a costof-living adjustment for hospitals in Alaska and Hawaii, and for plant/fixed capital costs, an area construction cost adjustment).
- We stated that we would determine the hospital-specific portion of the capital payment on the basis of each hospital's allowable capital costs for plant and fixed equipment and for moveable equipment in each year of the transition periods that we proposed for each classification of equipment.
  - · We proposed to provide-
- A ten-year transition period for incorporating capital payments for plant and fixed equipment; and

- —A two-year transition period for incorporating capital payments for moveable equipment.
- We proposed to amend the existing payment policy for outliers (42 CFR Part 412, Subpart F), which is authorized by section 1886(d)(5)(A) of the Act, by adding a portion of the Federal capital

payment to the pool set aside for total outlier payments. We proposed that the cost outlier policy (§ 412.84) would be based on inpatient operating costs including capital, and that we would pay cost outliers only when inpatient operating costs (including capital) for a case are above the cost outlier threshold.

 We proposed that the blending of Federal and hospital-specific portions of the capital payment for plant and fixed equipment during the transition period would be weighted heavily toward the hospital-specific portion in the earlier transition years as follows:

minute the production of the state of the st	Plant and fixed equipment		Moveable equipment	
	eduib	ment		The Late
Cost reporting period beginning on or after	Federal (percent)	Hospital- specific (percent)	Federal (percent)	Hospital- specific (percent)
scal year:		d'anilor	12 10 0	preda
1988	5	95	33	67
1989	10	90	67	3
1990	15	85	100	
1991	20	80		
1992	25	75		
1993	30	70		200000000000000000000000000000000000000
1994	40	60		
1995	50	50		
1996	22	35		
1997	80	20		
1998	100	0		The second second

- We stated that we would determine the hospital-specific portion based on Medicare's share of total allowable capital costs of plant and fixed equipment, and Medicare's share of total allowable capital costs of moveable equipment (rolling base) subject to the applicable blending percentages in each year of the transition for each hospital.
- Puerto Rico hospitals would be included in the prospective capital payment process in accordance with sections 1886(d)(9) and (g)(3)(A) of the Act.
- For FYs 1988 and 1989, we proposed to adjust the capital payment amounts (Federal and hospital-specific portions) in order that the aggregate capital payment amounts under the prospective payment system approximate the aggregate capital payment amounts that would have been made, taking into account the reductions prescribed under section 1886(g)(3) of the Act, on a reasonable cost basis during FYs 1988 and 1989.
- We proposed to make capital payments to new hospitals on the same basis as all other hospitals subject to the prospective payment system, using the rolling base approach and the applicable Federal/hospital-specific blend for the Federal fiscal year in which the hospital first participates in the Medicare program.
- We proposed to exclude sole community hospitals from inclusion of

capital into the prospective payment system for cost reporting periods beginning before Ocother 1, 1990, in accordance with section 1886(g)(3)(C)(i) of the Act.

 We stated our belief that the proposed capital payment policy would negate the need for a further distinct capital exceptions process.

We think it is important to note that, in developing the capital payment policy described in the May 1987 NPRM, we took into consideration ProPAC's 1986 recommendations concerning capital payment policy and numerous public comments on the June 3, 1986 NPRM (51 FR 19970) in which we initially proposed to incorporate capital costs into the prospective payment system. Based on those recommendations and comments, we significantly revised the original proposal, published in the June 1986 NPRM, in the following ways before publishing the May 1987 NPRM:

- We separated capital costs for plant/fixed equipment from moveable equipment.
- We extended the transition period from four years to ten years for plant/ fixed equipment.
- We elected to use a rolling base rather than a fixed base year for hospital-specific costs.
- We elected to update the standardized average capital costs through FY 1990 by the estimated actual increase in capital costs per case rather

than the overall prospective payment update factor.

In response to comments we received on the June 3, 1986 and May 19, 1987 proposed rules, we are incorporating capital costs into the prospective payment system as we had proposed in the May 19, 1987 NPRM subject to the following modifications (which are more fully discussed in section II of the preamble below):

 We are revising the length and blend of the transition period for incorporating capital costs for moveable equipment into the prospective payment system pursuant to the following schedule:

Cost reporting	Moveable equipment		
period beginning on or after fiscal year	ter fiscal Federal		
1988	5	95	
1989	10	90	
1990	15	85	
1991	20	80	
1992	30	70	
1993	50	50	
1994	75	25	
1995	100	0	

 We will make additional capital payments to hospitals that are financially disadvantaged during the capital payment transition period by the changeover from cost reimbursement to prospective payments for capital. Under this provision, an exceptions process is established pursuant to the general authority granted to the Secretary under section 1886(d)(5)(C)(iii) of the Act to provide for exceptions and adjustments to prospective payment amounts as is deemed appropriate. The amounts to be paid under this exception process will be obtained by reducing the average standardized capital payment rates by five percent of the total Federal capital payments.

The criterion we apply to determine the eligibility of a hospital for a payment in addition to its prospective capital payment amount is whether the portions (for plant/fixed equipment or moveable equipment, or both) of a hospital's actual, allowable inpatient capital costs not paid through the hospital-specific portions (after applying the mandated reductions under section 9303 of Pub. L. 99-509 of seven and ten percent for FYs 1988 and 1989, respectively), are 175 percent or more than the total of the hospital's Federal capital payments (excluding payments for the hospitalspecific portion) for that period. For FYs 1995-1997, the exceptions policy will only apply to plant/fixed equipment since moveable equipment will be paid based on 100 percent of the prospective moveable equipment rate.

The amount of the additional capital payment for hospitals meeting the eligibility criterion will be equal to 70 percent of the difference between 175 percent of the hospital's total Federal capital payments (excluding payments for the hospital-specific portion) during the transition period and its portion of the actual allowable inpatient capital cost not paid through the hospitalspecific portion, as established by the cost report for the applicable period. We note that the Federal capital payments do not include the hospital-specific portion of a hospital's total capital payment. A five percent pool has been established for payment of the adjustment, and we will announce prior to October 1 of each year of the transition any changes to the threshold level, if needed, to maintain the five percent pool. These additional capital payments will be adjusted retroactively for each cost reporting period during the transition based on changes in each hospital's actual, allowable inpatient capital cost as determined in its Notice of Amount of Program Reimbursement under cost reimbursement principles pursuant to section 1861(v) of the Act and implementing regulations at 42 CFR Part 413, Subpart G, §§ 413.130 through 413.154, and 42 CFR Part 412, Subpart D. §§ 412.65 through 412.68.

We are providing, below, two examples to demonstrate the exception payment process for a hospital that is financially disadvantaged during the transition period under the prospective capital payment system. Example 1 is applicable for FY 1988, a transition year in which the blended percentages for plant/fixed equipment and moveable equipment are the same. Example 2 is applicable for FY 1992, a transition year in which the blended percentages for plant/fixed equipment and moveable equipment are different.

Example 1-Hospital A submits its cost report for FY 1988 to its fiscal intermediary showing the following

capital data:

· Medicare allowable capital cost of \$1,000,000 (\$800,000 for plant/fixed equipment and \$200,000 for moveable equipment).

· A hospital-specific portion of Medicare allowable capital cost of \$883,500 [(\$800,000 × .93 (reduction under section 1886(g)(3) × .95 (hospitalspecific blend)) plus (\$200,000 × .93 ×

· Total capital payments received from Medicare during the fiscal period of \$898,500 [(\$883,500, the payment for the hospital-specific portion) plus (\$15,000, the amount received under prospective capital payment rates which includes payments for plant/fixed equipment, moveable equipment, outliers, indirect medical education cost adjustment and disproportionate share adjustment)].

Hospital A has requested additional payment (an exception) for capital costs pursuant to § 412.68. After a review of the data submitted, the fiscal intermediary has determined that Hospital A is due an additional \$14,175 pursuant to § 412.68. The fiscal intermediary computed the \$14,175 based on the following steps:

(1) Medicare capital cost not paid as part of the hospital-specific portion. Note that the hospital-specific portion is 95% for FY 1988.

 $\$1.000.000 \times .93^{1} \times .05 = \$46,500$ 

(2) Payments received under the prospective capital payment rates times the exception threshold pursuant to § 412.68.

 $$15,000 \times 1.75 = $26,250$ 

(3) Exception amount determined based on 70% of the difference between steps 1 and 2.

 $(\$46,500 - 26,250) \times .70 = \$14,175$ 

Example 2-Hospital A submits its cost report for FY 1992 to its fiscal

intermediary showing the following capital data:

· Medicare allowable capital cost of \$1,000,000 (\$800,000 for plant/fixed equipment and \$200,000 for moveable equipment).

· A hospital-specific portion of allowable plant/fixed equipment of \$600,000 (\$800,000 times 75% fixed hospital-specific blend) and a hospitalspecific portion of moveable equipment of \$140,000 (\$200,000 times 70% moveable hospital-specific blend).

 Total capital payments received from Medicare during the fiscal period of \$865,000 [[\$740,000 for the hospitalspecific portion) plus (\$125,000 for the prospective capital payments for plant/ fixed and moveable equipment)].

Hospital A has requested an additional payment (an exception) for capital costs pursuant to § 412.68. After a review of the data submitted, the fiscal intermediary has determined that the hospital is due additional payments of \$28,875 pursuant to § 412.68. The fiscal intermediary computed the \$28,875 based on the following steps:

(1) Medicare plant/fixed equipment cost not paid as part of the hospitalspecific portion. Note that the hospitalspecific portion is 75% for FY 1992; 25%

is not paid.

\$800,000 × .25 = \$200,000

(2) Medicare moveable equipment cost not paid as a part of the hospitalspecific portion. Note that the hospitalspecific portion is 70% for FY 1992; 30% is not paid.

\$200,000 × .30 = \$60,000

(3) Total Medicare plant/fixed and moveable equipment costs not paid as part of the hospital-specific portion in FY 1992. Add the values from step 1 and step 2.

\$200,000 + \$60,000 = \$260,000

(4) Payments received under prospective capital payment rates for plant/fixed and moveable equipment times the exceptions threshold pursuant to § 412.68.

\$125,000×1.75=\$218,750

(5) Exception amount determined based on 70% of the difference between step 3 and step 4.

 $(\$260,000 - \$218,750) \times .70 = \$28,875$ 

Note: Since reductions under section 1886(g)(3) of the Act do not extend beyond FY 1989, no reduction is reflected in this example. Also, the FY 1988 thresholds were used in this example, since we do not yet know the FY 1992 thresholds.

· The construction cost index to be applied to the Federal capital rate for

¹ Capital reductions for FY 1988 under section 1880(g)(3) of the Act.

plant/fixed equipment in Puerto Rico is 1.000. We do not have adequate data on which to base a Puerto Rico-specific ratio or proxy. We do not believe any adjustment relative to other areas would be appropriate at this time since the index will be applied only to the 25 percent national portion of its payment.

#### II. Comments and Responses

A total of 100 sets of comments concerning the May 1987 proposed rule were received timely. The commenters included 58 individual hospitals, ten hospital systems or corporations, 10 local or State hospital associations. three individuals, one U.S. Senator, 11 national health care assocaitions, two Medicare fiscal intermediaries, ProPAC, two law firms, and an insurance company. All of the commenters except for ProPAC voiced opposition to the incorporation of capital costs into the prospective payment system, both in general (by suggesting for example, that further study of the subject is necessary and requesting that we delay the incorporation) and with respect to specific issues. However, some commenters supported specific provisions of the NPRM, such as standardization for indirect medical education and the capital expenditure agreements policy. A few commenters, primary hospital associations, generally endorsed use of a construction index. Among the many issues addressed in the proposed rule, the following subjects received the majority of comments:

Adequacy of payment rates.
Split between plant/fixed equipment and moveable equipment.

· Construction cost index.

· Update factor.

· Urban versus rural rates.

· Outliers.

· Exceptions policy.

The contents of the proposed rule, the public comments, and our responses to the comments are discussed below.

We are responding to two general comments here rather than in one of the more issue-specific areas below.

Comment: The overwhelming majority of commenters opposed incorporating capital costs into the prospective payment system. The commenters asserted the following:

· The payment levels were grossly

inadequate.

 Hospital capital cost cycles are— Unevenly distributed over time;
 Composed generally of fixed costs;
 and

-Not amenable to substitution.

Thus, the incorporation of capital costs into the propective payment system would result in significant over- and underpayments (that is, a

maldistribution of payments) for many hospitals at the end/beginning of a capital expenditure cycle.

The capital data were flawed or inadequate.

 The proposed rule did not account for significant reductions in occupancy rates subsequent to FY 1984 (the base year for establishing the Federal capital rates).

 No additional savings would be generated due to the capital reductions under section 9303 of Pub. L. 99–509.

 Reasonable cost reimbursement rules took into consideration hospital

capital cost cycles.

Response: We acknowledge that in moving to an average pricing system to establish payments for capital expenditures for inpatient hospital services, we will not recognize the individual hospital capital cycle experience in the standardized portion of the prospective capital payment, and that the capital rate will be lower than a hospital's capital cost per case in roughly half of the cases. However, we believe these problems are alleviated, to some extent, by the provisions to have an extended transition period during which hospitals will receive a hospitalspecific portion of the prospective capital payment based on their actual allowable capital costs and the transition blend, which is heavily weighted toward the hospital-specific portion for several years. We believe that those hospitals with substantially higher capital costs per discharge than the Federal portion of the capital payment will have adequate time under the transition period to adjust their operations and financing to meet the relatively lower payment levels by the time the Federal rate becomes a major portion of their capital payment. We are also making changes in this final rule based on these comments to aid hospitals that are unable to adjust their capital expenditures timely. However, because the Federal capital payment is based on industry-wide averages, we expect that the capital payment for most cases will be appropriate for the majority of hospitals.

While specific data concerns are addressed below, we wish to point out that with respect to setting the Federal capital payment rate, the most recent, audited cost report information available to us was used to set the rates. We believe those data are valid and reliable since they are based on the reports of the nearly complete universe of hospitals subject to these rules. All of the actual hospital capital costs reported were paid for on that basis for that period. Thus, we do not believe the cost per case used was understated. Further,

since actual industry inflation rates (based on data provided by the American Hospital Association) were used to update the FY 1984 base year costs per case to FY 1988 levels (modified by legislative reductions imposed for FYs 1987 and 1988 under section 9303 of Pub. L. 99–509), the Federal rates are representative of the actual national cost per discharge levels that hospitals can expect to experience.

With respect to occupancy rate declines experienced by hospitals, these declines are taken into account in the capital cost per case projections that were used to determine the FY 1988 prospective capital payment rates. Such changes are taken into consideration automatically in calculating the projections, since we allocate the aggregate of costs in a period to Medicare's share of costs on a per admission basis for each such period. Thus, the cost per case inflation rate from year to year is dependent on changes in discharges due to occupancy level changes.

We also wish to clarify that this initiative is not intrinsically designed to increase Medicare program savings, but rather to correct the lack of incentive for controlling capital expenditures under cost reimbursement rules. Since no substantive alternative to continued cost reimbursement was suggested by commenters on either of the capital payment proposed rules, we do not believe that a delay in implementing this change would be appropriate in light of—

- Current Medicare program objectives to make hospitals more efficient;
- The recommendations of ProPAC;
  and
- The discretionary authority granted to the Secretary under the statute on this matter.

Comment: Many commenters pointed out that the average pricing method of making capital payments would disadvantage certain categories of hospitals with greater than average capital costs such as—

- Tertiary care hospitals which place heavy reliance on specialized, moveable, diagnostic and care equipment;
- Teaching hospitals with similar problems; and
- Rural hospitals, which maintain that the average age of their plants and equipment is greater than urban hospitals and that their lower rates for capital payments will exacerbate the problems they believe result from operating payment rates much lower than the rates for urban hospitals.

Further, the commenters pointed out that the impact on such hospitals due to conformance of State Medicaid plans to Medicare capital payment rules will accentuate financial results of this major program revision.

Response: We believe that the average pricing method for capital payments is the best approach to use for structuring prospective capital payments for hospitals. Like the method used to establish other prospective payment operating rates, this method does not differentiate between hospitals on the basis of size or efficiency. Since section 1886(d) of the Act provides for adjusting prospective capital payments for casemix complexity, disproportionate share, indirect medical education, and referral center considerations, capital payment levels for those categories are expected to be responsive to such hospitals unique capital needs in the same manner as prospective payments for other inpatient operating costs.

With respect to rural hospitals in general, we point out elsewhere in this issue of the Federal Register that we are required to compute rates under the prospective payment system on the basis of urban and rural averages, pursuant to sections 1886(d)(2)(D) and (d)(3)(D) of the Act. All data available to us regarding urban and rural costs for both operating and capital expenditures reflect the distinction between the urban and rural rates under the Medicare program. On that basis, we do not believe that the capital payment rates vill selectively disadvantage rural hospitals.

We do not expect that reimbursement changes by State Medicaid programs will be any more significant than the comparable changes resulting from implementation of prospective payments for other inpatient hospital operating costs occurring after October 1, 1983 (the inception of the prospective payment system).

#### A. Adequacy of Payment Amounts

#### 1. Base-year data

We proposed to use inpatient capital cost data from FY 1984 Medicare cost reports (latest available audited data) in determining the Federal capital payment rates.

Comment: Several commenters
expressed concern about the adequacy
of the data used to establish the capital
payment rates. They cited problems
with the adequacy and accuracy of the
cost report data used to calculate the
capital rates including—

Accurate classification of capital costs;

- Inadequate data base for all hospitals and units;
  - · Missing and erroneous data entries;
- Lack of a complete description of computation methods; and
- Unsettled cost reports for the base year.

Response: As we noted above, we are using the most recent audited data available from a statistically reliable volume of hospitals to determine the Federal capital payment rates. That data source is the set of Federal FY 1984 hospital cost reports as received. audited and submitted by Medicare fiscal intermediaries through our Hospital Cost Report Information System (HCRIS). More than 90 percent of all hospital cost reports form the basis for this data base which is used to set the prospective capital payment rates. This level of response permits us to conduct a statistically reliable analysis using HCRIS. Also, the data used were provided by hospitals that were reimbursed for their capital costs on the basis of the cost report information they furnished, and we, therefore, have every expectation that the same information is a valid basis on which to establish the average capital cost per discharge under this final rule. Although some cost reports for the period are not settled, all were received, and the vast majority audited by the fiscal intermediaries.

We agree that insufficient data were available at the time the May 1987 NPRM was issued that would support the separation of directly assigned capital costs into its various component costs. However, we did conduct a survey of a sample group of hospitals and obtained these component costs. The final rule includes those data which permit the analysis of the fixed/ moveable split of directly assigned capital costs as described in the response below regarding this separation. Thus, we believe that the data and procedures we used to calculate the prospective capital payment rates are adequate and accurate.

#### 2. Updating

We proposed to update hospitals' FY 1984 costs per case through FY 1987 by the estimated actual increase in capital costs per case for purposes of establishing the rates. We stated we would also update costs for FYs 1988 and 1989, subject to the reductions under section 1886(g)(3) of the Act, under reasonable cost principles. For FY 1990 onward, the update provisions under section 1886 (b)(3)(B) and (e)(4) of the Act will be applied to the capital

rates as to all other inpatient hospital operating rates.

Comment: Many commenters expressed their confusion and apprehension about the update factors and the mechanism for prospective capital payments. They cited a lack of information in the proposed rule regarding the factors used to trend forward the FY 1984 base year cost per case to the first transition year (FY 1988). The commenters said that there is no indication of whether or how the prospective payment update factor will be adjusted to ensure that capital cost changes are properly reflected in future updates. The commenters also cited the need to disclose how the capital market basket will be constructed and applied, considering such factors as the volatility of interest rates, moving averages for the useful life of assets, new technology, and other factors. They were concerned that updates in the future will be used to cut back Medicare expenditures, ignoring the long range impact on hospitals and, thus, violating the requirements of Executive Order 12291 regarding disclosure of the long range impact in rulemaking.

One commenter stated that the update for capital payment rates should be established independent of the update for other prospective payment system operating rates during the transition period. Several commenters also stated that the inflation of the cost per case from FY 1984 to FY 1988 does not properly recognize technological changes and occupancy rate changes. Finally, one commenter suggested that the capital payment reductions contained in section 9303 of Pub. L. 99-509 should not be continued in the base to which update factors are applied after Federal FY 1989, and that the final rule on prospective capital payments should address this matter specifically.

Response: We agree that we neglected to indicate the amounts and source of update factors used to trend the average cost per discharge in the capital payment standardized rate base year (FY 1984) through the first two transition years, which are required to be budget neutral under section 9303 of Pub. L. 99–509 (FYs 1988 and 1989). The amounts used were the estimates of actual industry-wide capital expenditure increases based on the American Hospital Association panel survey data and converted to a per admission basis. The update factors for each year are:

FY 1985—17.29 percent FY 1986—10.86 percent FY 1987—7.06 percent FY 1988—6.83 percent FY 1989—5.86 percent

However, we believe that the budget neutrality requirement methodology and its impact on the calculation of updates for FYs 1988 and 1989 were thoroughly explained. Of course, that requirement, along with adjustments required to standardize the Federal rates, appears to reduce the average increase for inflation over this period as noted by several commenters. Nonetheless, no other action or adjustment to the rates was made to reduce them or to increase program savings. Further, the update to the rates between the base year (FY 1984) and FY 1988 takes into account hospital practice patterns and effective technologies. Thus, the resulting rates reflect our best estimate of the Medicare inpatient capital payments that would be made in FY 1988 subject to the per case reductions required under section 9303 of Pub. L. 99-509. We will revise the hospital market basket to reflect capital items and services, as discussed below. Therefore, updates to the Federal capital payments made after FY 1989 will be included in the overall update to the prospective payment system in order to fully integrate capital payments into the prospective payment system.

With respect to the concern for the reduction (pursuant to section 1886(g)(3) of the Act, as added by section 9303 of Pub. L. 99-509) in the base used to set capital payment rates after FY 1989, we believe that we have no discretion but to use the allowable cost base, as reduced under section 1886(g)(3) of the Act, in trending forward the base year amounts in setting the prospective capital rates. Since the years in question (FYs 1987 through 1989) are subject to statutorily mandated reductions, those amounts must be used as the basis for capital payment rate setting in the same manner as the rate of increase limits. established under section 1886(b) of the Act, were incorporated for other inpatient operating costs in setting the initial prospective payment system

#### 3. Urban Versus Rural Payments

We proposed to compute average standardized rates for plant/fixed equipment and moveable equipment, for all urban hospitals and for all rural hospitals in the United States and the District of Columbia, and for urban and rural hospitals in Puerto Rico. The national urban and rural averages would be discharge weighted in the same manner as other prospective payment rates.

We refer the reader to the general comments and responses discussed above.

#### B. Standardizations

#### 1. Construction Costs

We proposed to standardize plant/ fixed equipment capital costs for area construction costs using the Dodge/Data Resources, Incorporated (DRI) Construction Potentials database as the source for developing a construction costs index. We still believe that capital payments should vary geographically by construction costs.

Comment: Many commenters pointed out specific locations in which the construction cost index varies significantly across geographic boundaries, and stated they did not see why such variations should occur.

Response: There are a number of reasons why the construction cost index varies across local geographic boundaries, such as differences in the wage rates of construction workers or differences in the cost of materials used in construction. We expect local variations in the cost of construction because of the degree of competition (or lack thereof) between construction companies and the general economic

conditions of an area.

As explained in the NPRM, there was considerable year to year variation in the relative cost of construction in many metropolitan statistical areas (MSAs)/ New England County metropolitan areas (NECMAs). We believe it was important to average over an extended period (15 years) to smooth these fluctuations. By averaging over 15 years, the total volume of construction used to compute the index for each MSA/NECMA is increased, which helps prevent a single very expensive, low volume project from unduly influencing the index for an MSA/NECMA. If the cost of construction in an area has changed significantly relative to the national average during the past 15 years, it is possible this longer averaging period could cause an MSA/NECMA with the majority of the construction in its index in the past few years to have a different index than an adjacent area with little recent construction to influence its index. Even though the averaging technique may create a few boundary problems, we believe it is important to have a construction cost index that does not have wide fluctuations over short periods of time. Therefore, we believe this approach and the data for establishing the index are the best available way to measure geographic variation in hospital construction cost. We will continue to refine this index to measure local variation more precisely.

Comment: Several commenters provided specific local construction or cost of living data that they believe

showed the construction cost index did not sufficiently compensate their area. relative to adjacent areas. Several suggested either the data must be flawed, or HCFA's manipulation of the data was incorrect.

Response: Data for this index were chosen from among a number of alternatives, as discussed in the May 1987 NPRM, because we believed it to be the best overall source for developing a prospective payment construction cost index (52 FR 18846). In some areas it may be possible to cite specific local data, such as wages for construction workers typically employed in hospital construction, which would appear to indicate the indices in adjacent areas should not vary. However, as we stated in the NPRM, there are significant fluctuations in the yearly construction cost data. Because of these fluctuations. we do not believe that a reliable index can be developed from one year of data, nor could we make appropriate comparisons from one year of data. We have not found evidence to suggest an index computed from an alternate data source would be more accurate nationally than the current index. We do not believe the data are flawed or that our methods for computing the index are inappropriate or incorrect.

Comment: A number of commenters stated that the construction cost index is invalid because it is based on data that include the construction costs of schools, libraries, churches, and other public buildings that do not reflect the higher and very different construction costs incurred in hospital construction. The increased costs associated with hospital construction (such as, the heavy-duty electrical system or built-in gas systems) generally arise from the life and safety standards required for hospital care. One commenter noted that, in particular, for those areas in which there are sufficient hospital construction cost data, the construction cost index should not include nonhospital construction costs.

Response: The construction cost index measures the relative costs of construction in an area, not the absolute costs. We believe that the types of construction (hospital and nonhospital) used to compute the construction cost index represent the same market for contractors as hospitals, and vary in construction costs in a manner similar to hospitals. As stated in the NPRM, we found using hospital-specific data results in a number of MSA areas having five or less years of data to average during the past 15 years. An index computed using hospital specific data from DRI has a 31 percent larger

standard deviation and a 38 percent wider range. Based on these results, we do not believe an MSA level index based on hospital only construction would be a better indicator of geographic variation in hospital construction cost than the index given in the NPRM. However, in the future, we may phase out nonhospital construction as we obtain more data.

Comment: Many commenters suggested that an index based on historical average construction costs is inappropriate because it fails to account for where a hospital is in its construction/depreciation cycle.

Response: We disagree with the commenters. The purpose of the construction cost index is to measure differences in the costs of construction among geographic areas. The index is not intended to adjust for differences in the construction/depreciation cycle (age) of a hospital's fixed capital.

Comment: Several commenters suggested that the construction cost index should be based on replacement capital costs rather than historical capital costs because use of historical capital costs fails to account for the financial position of a hospital in its construction/depreciation cycle. In addition, the construction cost index should account for the higher capital construction costs associated with new technology.

Response: Historical construction cost data are used because we believe they are the best data available to measure the geographic variation in construction cost. Since we do not have a reliable way for projecting replacement construction cost, we are unable to establish an index based on replacement capital cost. However, since we use a 15 year average to determine each MSA's/NECMA's construction index, future replacement costs, which include new technologies, will be represented in the construction cost index in future updates.

Comment: Several commenters suggested that the construction cost index should be computed from hospital specific data for those areas that had a large volume of hospital construction activity, and an alternative approach should be developed for those areas in which sufficient hospital-specific data do not exist. One approach would be to use the hospital-specific index of a similar geographic area. Another suggestion was to conduct a survey of general contractors in those areas for which additional data were needed to estimate local hospital construction costs.

Response: In developing the construction cost index, we considered

a composite index based on hospital only data for those areas that exceeded a certain threshold of hospital construction, with an alternative supplemental data source for those areas in which there was insufficient hospital construction to develop a reliable index. A preliminary analysis of our data suggested the areas with sufficient hospital data to develop a reliable index would be the 50 or so largest MSAs, and many State rural areas. However, we were concerned that developing an alternative data source for the remaining perhaps twothirds of the areas could bias the index. We therefore decided not to use this approach but to use a construction cost index in which all areas of the country are treated in exactly the same manner.

Comment: One commenter suggested that construction costs do not vary as much at the MSA level as the construction cost index indicates, and a regional index would be more appropriate.

Response: We acknowledge that whenever geographic boundaries are established, boundary problems may occur. We recognize that a regional construction cost index would reduce the number of boundaries; however, such an index would not recognize local variations in construction costs. Thus, we believe that an index based on the MSA/NECMA level is the most appropriate measure of fixed capital cost variation.

Comment: Several commenters stated that the construction cost index is biased toward urban hospitals, even though rural hospitals typically use contractors from urban areas who charge the same or more (because of transportation costs) for rural construction as for urban.

Response: We believe that a portion of the difference in construction costs between regional and urban areas results from the nature of urban construction, which tends to be more vertical (that is, taller buildings) than in rural areas because of the higher premium on space. In addition, the unweighted average difference between the construction cost indices for urban and rural areas is not large, with the average for all urban areas being 1.008. and .948 for all rural areas. This point is further exemplified by the construction cost indices for the western mountain States, in which rural transportation costs are probably quite significant. For these States, the unweighted average urban index is .928, while the average rural index is .934.

Comment: Two commenters alleged that because the construction cost index

has an overall east coast and metropolitan bias, it is inappropriate.

Response: We agree that the construction cost index tends to be higher in the eastern part of the country than in the western part of the country, but we do not believe there is any systematic bias in the data. Both the hospital specific data and the data for other institutional construction included in the index separately exhibit this tendency, and we believe they are measuring appropriate construction cost differences in different regions of the country.

Comment: In response to our request in the NPRM for information and suggestions on how to establish a construction cost index for Puerto Rico, one commenter suggested direct collection of construction cost records and information from Puerto Rico hospitals.

Response: Prior to publication of the NPRM, we initiated telephone and written requests to obtain such hospital specific data from Puerto Rico, but have not yet received any data that could be used to develop a construction cost index for Puerto Rico. Because of our lack of Puerto Rico-specific data, and our analysis of the several proxies for Puerto Rico that we mentioned in the NPRM, we do not believe that we have sufficient information at this time to determine an appropriate index for Puerto Rico. We therefore have decided to use an index of 1.000, until a more appropriate index can be developed. This index is applied only to the 25 percent portion of Puerto Rico's Federal capital payment rates, which is based on the national Federal capital payment rates. The Puerto Rico-specific capital rates are not subject to this index.

Comment: One commenter suggested that the construction cost index is being improperly applied to all fixed capital costs (which include depreciation, lease/rental, interest, insurance and taxes), while it should be applied only to depreciation. Another commenter suggested that the index should take into consideration "soft" construction costs (that is, architect's fees, legal fees, accounting fees, finance costs during the construction period, and development overhead costs), which the commenter estimated to represent between 15 and 35 percent of total construction cost.

Response: Although the construction cost index only measures specifically the cost of constructing a fixed asset, we believe that, in general, the interest resulting from a hospital's debt along with insurance and tax costs vary in direct proportion to the cost of the fixed asset constructed. We also believe the

geographic variation in construction costs represented by the index is a reasonable proxy for geographic variation in the soft construction costs alluded to by the second commenter. Thus, we believe it is appropriate to apply the construction cost index to all of the fixed capital costs referenced by the commenters.

Comment: One commenter stated that because the construction cost index does not take into account inflation, many hospitals located in the same geographic area to which the same index is applied, will either be undercompensated or overcompensated for their capital construction costs depending on the year in which they incurred their construction costs.

Response: The construction cost index was designed to measure the relative differences in the costs of construction between geographic areas for assets purchased in the same year. The index, which is based on a 15 year average, is calculated in such a way as to remove the effects of construction cost inflation over time. Since the index is not dependent on the age of the asset, it does not result in undercompensation or overcompensation to hospitals that have different fixed asset age values.

Comment: Several commenters requested that the construction index should be updated on an annual basis to reflect actual construction costs.

Response: We agree that the construction cost index should be reviewed on an annual basis, and we will update the published values of the index as appropriate.

Comment: One commenter recommended using cost report data to develop a construction cost index.

Response: While cost report data are a basis upon which an index could be developed, it would be necessary to have several years of reliable data to smooth out yearly fluctuations. Currently, we do not have the necessary cost report data to perform the appropriate analyses for determining whether a geographically adjusted construction cost index could be developed from cost report data. Also, it would be difficult to establish a common unit for measuring relative cost, since depreciation per square foot would need to be standardized by the age of the asset being depreciated.

Comment: One commenter suggested that because many of the reported values were excluded from the construction cost index calculations, it might be flawed.

Response: Out of 365 MSA/NECMA and State rural areas, each with 15 years of data, 15 of the (365 x 15) data "cells" were excluded as outliers. This results in less than .28 percent of the data being excluded in calculating the index, which we believe is not significant. Further, we believe it is appropriate to exclude this small number of outliers from the construction cost index.

Comment: Two commenters requested that an appeals process be established for hospitals to request reconsideration of their assigned construction cost index

Response: Because of the complex multi-year formula we used to establish the national hospital construction cost index for 365 MSAs/NECMAs, we do not believe that an appeals process would be the most appropriate mechanism for addressing construction cost values that hospitals believe are aberrant. If hospitals have problems with their construction cost index value, we encourage them to submit data or an alternative methodology that they believe will improve the national construction cost index.

Comment: One commenter stated that central city construction costs were considerably higher than suburban construction costs, and that the construction cost index should address those cost differences.

Response: To develop an index that distinguishes between central cities and suburban ring areas within an MSA/NECMA, it would be necessary to develop geographic definitions to properly distinguish between central cities and suburban ring areas. We are not aware of a way to make such distinctions uniformly, but would consider use of such definitions if they become available in the future.

Comment: Two commenters questioned the disparity between their area wage index values and their construction cost index values in light of the fact that labor constitutes 60 percent of construction costs.

Response: While there is a positive correlation between the index for construction costs and the index for average wages of hospital workers, the correlation is not high. There are a number of factors that could cause the construction cost index to vary geographically from the wage index, including weather, terrain, and cost and proximity of building materials. Finally, the construction cost index is a multiyear average index, while the wage index is based on a single year of data. In view of these factors, we do not believe that an implication arises that the construction cost index is incorrect if it is not similar to the wage index.

#### 2. Indirect medical education

We proposed to standardize capital plant/fixed equipment and moveable

equipment costs for indirect medical education, case mix, disproportionate share of low-income patients, and other factors.

Comment: The majority of comments received about our request for evaluation of whether to standardize the prospective capital payment rates for indirect medical education and disproportionate share of low-income patients were in favor of such adjustments to the rates. Several commenters objected to standardization for these items observing that it is not clearly established by studies that capital costs are affected in the same manner that operating costs would be in these situations. In addition, commenters also expressed concern whether case mix should be used to standardize costs due to the methods used to allocate capital costs to various portions and departments of hospitals (square foot basis rather than department of major use), which allegedly distorts the resulting diagnosis-related group (DRG) weights, and, thereby, the case mix, with respect to capital item use intensity.

Response: The few comments we received that objected to standardization of the Federal rates for disproportionate share of low-income patients, indirect medical education, and other factors, were usually related to the fact that such standardization reduces the capital rates somewhat. However, we believe that this redistribution of payments is required under the statute since the removal of any statutory distinction between inpatient capital and other inpatient operating costs subjects all rates computed under the prospective payment system to the same standardization requirements.

With respect to the appropriateness of standardization by case mix in the capital payment calculation, we believe that standardization is required by section 1886(d)(2)(C)(iii) of the Act, and application of the DRG weight by section 1886(d)(2)(G)(ii)(II) of the Act.

We also believe that by using the charge data to recalibrate the DRG weights, as described in the September 3, 1985 final rule (50 FR 35722), capital costs are reasonably represented in the relative weights used to compute the Federal capital payment amounts. Furthermore, as capital use intensity changes in an individual DRG, future recalibrations will take into consideration such changes and automatically adjust the payment levels.

C. Split of Plant/Fixed Equipment from Moveable Equipment

We proposed to split plant/fixed equipment costs from moveable equipment costs by developing, for each hospital, ratios of plant/fixed equipment costs to the total capital costs and of moveable equipment costs to the total capital costs.

Comment: More than one-third of the commenters objected to the proposed split and separate transition period of moveable equipment from plant/fixed equipment. They cited the following reasons for their opposition:

- The split would redistribute capital payments in a manner that would seriously disadvantage hospitals with high moveable equipment costs.
- Significant proportions of moveable equipment are financed through longterm debt arrangements, contrary to the assumption stated in the proposed rule.
- More than three-fourths of the major moveable equipment listed in the AHA useful life guidelines have useful lives of 10 years or longer, thus undermining any transition distinction rationale.
- Much moveable equipment cost is included in long-term debt costs obtained for plant/fixed equipment financing arrangements, thus skewing any cost report data used in determining the split.
- None of the proposed bases for allocating interest expense between moveable equipment and plant/fixed equipment can be expected to allocate properly between the two because each method will result in a bias in valuation toward one type of capital expenditure or the other.
- The policy ignores the wide variation in the proportion of actual cost distribution between the two categories among hospitals.

Response: The method used to determine the proportions of moveable equipment and plant/fixed equipment for purposes of setting capital payment transition rates was explained in the proposed rule (52 FR 18846). We believe that the cost report information on this split is generally reliable for hospitals that separately reported each category. and that no better source of direct data on this distinction has been developed. There is no evidence that hospitals grossly or consistently misreported the categories of capital expenditures identified on the Medicare cost reports. While reimbursement of capital costs was not dependent in any way on this designation of capital costs, there was no incentive for hospitals to misrepresent those costs.

Only for the cases in which hospitals directly assigned capital costs and did not separately assign these costs to moveable or plant/fixed equipment categories would distortions of the ratio between the two be reasonably questioned. However, as we noted in the proposed rule, a special data collection effort was undertaken to provide additional information on directly assigned capital costs, which disaggregated them between moveable equipment and plant/fixed equipment. This effort, using FY 1984 settled cost reports, provided a reliable sample for an analysis to determine whether our assumptions in making the split, as presented in the proposed rule, were supportable, or if an adjustment based on the special study should be made. Based on the results of that study, we find that the ratios developed under the assumptions in the proposed rule are to be adjusted by splitting any directly assigned capital costs reflected in a hospital's base year cost report between plant/fixed equipment and moveable equipment in the following manner:

Phenippi Milani Sakar Azzi	Urban (per- cent)	Rural (per- cent)
Ratio of Plant/Fixed Equipment Costs to Total Capital Costs	28.3	18.2
Total Capital Costs	71.7	81.8

Thus, we believe we now have an adequate and supportable basis on which to proceed with separate rates for each category.

While we recognize that each hospital's split of plant/fixed equipment from moveable equipment will vary from the proportions reflected in the capital payment rates, even significantly at times, the length and blending of the prospective payment transition period for the moveable equipment category is intended to moderate this impact, giving hospitals adequate time to adjust their situations Since the hospital-specific portion of the blend is important in ensuring that hospitals with widely divergent capital expenditure patterns are not adversely affected by the incorporation of capital payments into the prospective payment system, this concern of commenters added significantly to our decision to modify the transition rules (length and blend) for moveable equipment, as explained in our response to the comments regarding the length of the transition period below.

We believe that the modification to the transition period will correct the problems noted by commenters in regard to the longer term expenses encountered with major moveable

equipment. We also believe that distinguishing between moveable equipment and plant/fixed equipment under the refined cost finding guidelines, and the update of the list of plant/fixed equipment and moveable equipment, as discussed in the May 1987 NPRM (52 FR 18851-18855, and 18864), will properly identify the wide variations among hospitals in the proportion of debt between those categories allowing appropriate payment under the hospitalspecific portion of the capital payment blend for a longer period, so that no disadvantage will result to hospitals highly capitalized in moveable equipment.

Comment: Two commenters questioned whether the proposed regulations at § 412.67(d) that clarify cost finding distinctions in classifying plant/fixed equipment and moveable equipment, and the list of plant/fixed equipment and moveable equipment detailed in Appendix C of the NPRM, preclude the possibility of using other classification schemes or hospitals' past practices that may be appropriate as well.

Response: We believe that the approach for classifying plant/fixed equipment from moveable equipment, as stated in the May 1987 NPRM, is appropriate in terms of administrative efficiency. In addition, the list presented in Appendix C is based on usual industry practice and generally applied accounting procedures, as adapted from a comparable list of the American Hospital Association. However, we will provide a hospital an opportunity to appeal a classification if the hospital has had a longstanding past policy or practice of classifying items that is different from our approach, and that policy is not for the purpose of the hospital affecting its prospective capital payments. (We note that the use of a past practice of classifying equipment would not be appropriate, for example, if a building was classified as moveable equipment.) We are revising § 412.87(d) to implement this provision.

#### D. Length of Transition Periods

We proposed a ten-year transition period, heavily weighted toward the hospital-specific portion in the earlier transition years, for incorporating capital payments for plant/fixed equipment into the prospective payment system. Similar to ProPAC's recommendation, we proposed a two-year transition period for incorporating capital payments for moveable equipment into the prospective payment system.

Comment: A majority of the commenters not including ProPAC expressed opposition to the length of the transition periods. While the commenters were almost evenly divided as supporting or opposing the 10-year transition period for plant/fixed equipment, the vast majority objected to a two-year transition period for moveable equipment. In general, a number of commenters suggested that the moveable transition period should be between five and ten years. However, other commenters proposed that the transition period for moveable equipment should be the same as it is for plant/fixed equipment since it would take hospitals just as long to adjust to standardized payments for major moveable equipment. All those commenters who addressed the hospital-specific portion ("rolling base"). which is heavily weighted in the early transition year, were in favor of that

Response: Based on the commenters' suggestions regarding the inadequacy of the transition period and the blend for the movable equipment component of the prospective capital payment, and comments critical of the split between moveable equipment and plant/fixed equipment (discussed above), we are revising the moveable equipment transition and blend in this final rule. As shown below, the transition for moveable equipment is extended to seven years (a level between the fiveyear and ten-year levels suggested by commenters), and the blend of Federal rate and hospital-specific amount is made identical to the blend for plant/ fixed equipment in the first four years of the transition period.

Cost reporting	moveable equipment		
period beginning on or after fiscal year	Federal (percent)	Hospital- specific (percent)	
1988	5	95	
1989	10	90	
1990	15	85	
1991	20	80	
1992	30	70	
1993	50	50	
1994	75	25	
1995	100	0	

We believe that this change will ameliorate the problems cited by commenters on this matter. However, we do not believe that longer transitions for these capital components are essential to allow hospitals sufficient time to adjust their operations and financing to accommodate prospective Federal capital payment rates.

#### E. Exceptions Process

We proposed not to establish an exceptions process because we had adopted a number of options (such as, a longer transition period, and a "rolling" base year in determining the hospital-specific portion) that we believed negated the need for an exceptions process.

Comment: Nearly a third of the commenters objected to the lack of an exception process for hospitals that are financially disadvantaged by incorporating capital payments into the prospective payment system. All of the major hospital asociations opposed deferral of including an exceptions process for hospitals that are adversely affected by the capital payment methodology. They believe that the capital payment system will redistribute payments for capital items, often in ways that are not dependent on a hospital's efficiency, its position in the capital expenditure cycle and other critical factors. Many commenters indicated that even the longer transition with a blend heavily weighted toward the hospital-specific component in the early years would not be adequate to safeguard a critical shortfall of funds for some highly leveraged facilities.

Response: We recognize the concerns expressed over the lack of an exceptions process for hospitals that find themselves financially disadvantaged by the changeover to an average pricing system for capital expenditures. We are incorporating an exceptions process in this final rule, pursuant to authority granted to the Secretary under section 1886(d)(5)(C)(iii) of the Act, to deal with this problem. The exceptions process added here is based on that proposed as an option in the June 3, 1986 proposed rule (51 FR 19981), as modified in response to the comments received on that option. We had received 62 comments on that option, 18 opposed and 44 in favor. Of those commenters in favor, 25 suggested that less stringent or alternate criteria be used to establish eligibility for an exception payment adjustment to the Federal capital rates.

We are relaxing those criteria, as requested, to require that if the portion of a hospital's actual allowable inpatient capital cost not paid through the hospital-specific portion is 175 percent, or greater, than its Federal capital payments (based on the Federal prospective capital rates), the hospital will be eligible for an additional capital payment adjustment. We will pay an additional capital payment to recognize 70 percent of the difference between the portions (for plant/fixed equipment or movable equipment, or both) of the

hospital's actual, allowable capital cost not paid through the hospital-specific portions, and 175 percent of its Federal capital payments (based on the Federal prospective capital rates). A pool of funds is established for this additional payment process by reducing the average standardized capital payment rates by five percent of total Federal capital payments, based upon the comparable provision of the statute at section 1886(d)(2)(E) of the Act relating to outlier payments and the general intent of Congress and the Secretary to maintain payment equity in new prospective payment system program initiatives. We estimate that approximately 13 percent of prospective payment hospitals, nationally, will be eligible for this additional adjustment to their prospective capital payment. The 175 percent threshold level and the 70 percent payment factor are similar to the factors considered by Congress in 1986 during its deliberations on a capital exceptions policy. Each year we will evaluate the appropriateness of the threshold level. Unless we propose to do otherwise, the five percent additional payment pool and the 70 percent payment factor will be unchanged during the transition period.

#### F. Outlier Policy

We proposed that payment for captial day outliers be determined based on the same provisions in effect for noncapital day outliers. We proposed that payment for cost outliers be determined based on total inpatient operating costs including capital.

Comment: In response to both the June 1986 NPRM (51 FR 20028) and the May 1987 NPRM (52 FR 18850), many commenters objected to the policy proposed for making additional payments for day and cost outlier cases in accordance with current prospective payment system statutory and regulatory guidelines. Commenters cited the following objections to the capital payment outlier proposal:

 The capital payment pool should not be reduced by five percent to provide for capital outlier payment.

• A marginal cost factor rate of 60 percent should not be used; rather, it should be 80 percent as proposed for other inpatient operating rate outliers or 100 percent as hospitals now receive under cost reimbursement rules.

 The criteria and payment levels for capital outliers should be separated from those for operating rate outliers under the prospective payment system.

 Outlier payments should not be made for capital items and services at all because those are, by definition, fixed costs, whereas operating costs are marginal and variable and should be considered for marginal adjustment (outlier) payments in addition to a basic rate.

Response: There is no alternative, in our view, to the creation of a prospective capital payment outlier pool if such payments are to be made for capital items and services for cases meeting the day or cost outlier definitions contained in the prospective payment system statute. In that case, the provisions of section 1886(d)(2)(E) of the Act are clear in regard to creating a reserve of funds from the payment pool. As noted previously in these comments and responses, with the elimination of any statutory distinction between capital costs and other operating costs, effective with incorporation of the former in the prospective payment system per section 1886(a)(4) of the Act, the requirements for calculation and payment under section 1886(d) of the Act appear to be controlling. For this same reason, capital costs should be treated similarly with other inpatient operating costs, not as separate costs for outlier purposes.

We do agree that any increased marginal cost factor applied to other inpatient operating outlier payment situations would be applicable to capital outlier payment cases. However, since our intent is to replace capital payments under cost reimbursement rules with payment under the prospective payment

system, full allowable cost

reimbursement would be inappropriate.

The outlier methodology adopted in this final rule is generally the same as the methodology we published in the June 1986 NPRM. However, because section 206 of the Urgent Supplemental Appropriations Act for FY 1986 (Pub. L. 99–349) precluded the incorporation of capital payments into the prospective payment system for an additional year, we did not issue a final rule to do so at that time. However, we have considered the outlier comments on both the June 1986 and May 1987 NPRMs in the development of this final rule.

We are adopting, in general, the same methodology on payment of outliers as we had proposed in both NPRMs. That methodology combines the inpatient hospital operating and capital payment for outliers. The only methodological changes are necessitated by the features of the prospective capital payment system that are different from the June 1986 NPRM (such as, the construction cost index, separate plant/fixed and moveable payment rates, and different transition periods). These different features were stated in the May 1987 NPRM.

The methodology for computing outlier payments makes use of the labor-related, nonlabor related, and capital portions from the hospital market basket. These portions are derived from the same basic market basket data used in computing the prospective payment system update factor for FY 1987, as set forth in a September 3, 1986 final rule to update the prospective payment system (51 FR 31454), except that, for outlier purposes, the market basket includes a capital component that was described in the capital outlier methodology of the June 3, 1986 NPRM (51 FR 20028).

#### G. Interest Allocation

We proposed to allocate interest between plant/fixed equipment capital assets and moveable equipment assets by properly classifying capital interest expenses between plant/fixed equipment and moveable equipment.

Comment: Several commenters objected to any fixed formula approach to allocating interest expenses between moveable equipment and plant/fixed equipment. The comments generally indicated that Medicare cost report data were not precise enough to make an appropriate allocation of interest in all financing situations but that other allocation methods (such as historic book value or the first in/first assigned method) would significantly distort actual interest cost distribution to the degree that would seriously disadvantage hospitals financially. Two commenters suggested interest should be allocated to plant/fixed equipment

Response: With the changes we are making to the transition and blend for the moveable equipment component of prospective capital payments, we do not believe that any mechanism, other than Medicare's principles of reimbursement currently in effect, along with refinements noted in this document, and to cost reports to identify more precisely the cost of moveable equipment from plant/fixed equipment, will be needed to make appropriate capital payments for each category during the seven and tenyear transition periods for each component, respectively. Total payments for capital items and services of hospitals will not be affected by the split in the first four years of the transition period.

#### H. Puerto Rico

We proposed to establish separate adjusted average capital payment rates for urban and rural hospitals in Puerto Rico using FY 1984 Puerto Rico hospital data and, in general, the same methodology as that used for the fifty States and the District of Columbia,

Comment: One commenter noted that because the standardized Puerto Rico specific capital payment rate was very low, further audit of Puerto Rico hospitals' capital costs should be undertaken.

Response: We agree with the commenter. We have directly obtained, examined and extracted capital cost information from Puerto Rico hospitals' final, audited cost reports, and reviewed our rate calculation procedures. Those actions ensure that the revised standardized capital payment rates for Puerto Rico are accurate based on the data submitted by the hospitals and for which they have been reimbursed. In addition, since the blend for plant/fixed equipment and moveable equipment is the same for the first four years of the transition period, the specific capital payments for Puerto Rico hospitals is independent of the split between plant/ fixed equipment and moveable equipment capital rates during this time period.

#### I. Capital Expenditure Agreements

We proposed, based on our interpretation of section 1886(g)(3) of the Act, not to impose the requirements under section 1886(g)(1) of the Act which provide that if legislation concerning payments for inpatient hospital capital costs is not enacted before October 1, 1987, no payment may be made for capital costs of capital expenditures for inpatient hospital services in a State or jurisdiction if such expenditures are obligated after September 30, 1987. unless the State or jurisdiction has an agreement with the Secretary under section 1122(b) of the Act and under such agreement the State recommended approval of the capital expenditure.

Comment: Several commenters supported our finding that mandatory section 1122 agreements and review approval under section 1886(g)(1) of the Act were nullified by legislation that added section 1886(g)(3) of the Act. However, two commenters stated that further congressional confirmation would be appropriate.

Response: We believe that unless there is a further statutory change to the pertinent sections of the law, our understanding of section 1886(g)(3) of the Act is a reasonable and appropriate interpretation.

## J. Other

Comment: One commenter suggested that the recapture of depreciation rules at § 413.134 should be rescinded if the proposed rules are implemented due to reimbursement penalties against both

the seller and purchaser in such circumstances.

Response: The rules we are implementing to incorporate capital payments will not contravene those cost reimbursement rules applicable in determining the hospital-specific portion of prospective capital payments. Thus, purchasers in charge of hospital ownership situations will receive the same level of payment as the sellers were entitled to, subject to the capital payment transition blend and allowable cost determination rules.

While we agree that § 413.134 requires adjustments to the selling provider's allowable costs to recognize gains and losses on disposals of assets, we would not preclude payment of depreciation to the acquiring provider. In fact, in the proposed rule, we noted that assets may be revalued consistently with section 1861(v)(1) of the Act, and that revalued amount is used in determining the hospital specific portion of capital costs [52 FR 18853]. Therefore, we do not agree with the commenter that double penalties would occur.

Comment: Several comments were received regarding the inappropriate use of DRG weights in calculating any prospective capital payment amounts. Commenters allege that because different hospital departments tend to be involved with particular mixes of cases, many departments involved with treatment and care of patients are high capital item and service areas, but low on square foot area, and have a skewed impact on payments using operating cost-based weights and indices. This results from the fact that the weights and case-mix index are used in calculations for moveable equipment capital payment rates and plant/fixed equipment rates separately under the proposed rule. Thus, if a hospital is highly capitalized on major moveable equipment in ancillary departments and tends to treat patients in that setting, but the rates reflect a much lower moveable to plant/fixed equipment split ratio, the hospital would be disadvantaged. Other commenters note that on specific DRG weights, there is an irreconcilable loss in the average weights being used to pay on a case-by-case basis. The most blatant example is lithotripter treatment cases in which that DRG weight is half that of invasive techniques that require very limited capital intensive resources. Thus, hospitals would not recover appropriate costs for such cases until weighting averages change, which could be years away. Further, there is no indication our rules would ensure such technological changes will be made at all or in any reasonable timeframe to

assure adequate capital payment for hospitals that specialize in such activities.

Response: We refer the reader to the comment and response discussed above in section II.B.2. of the preamble concerning standardization. As we indicated above, use of total charges to recalibrate relative weights for intensity of resource use, including capital items and services increased by technology or other changes, will adjust the case-by-case payment levels.

Comment: Several commenters suggested that the mechanism for making blended capital payments during the transition period be specified in the final rule. Some suggested a standard schedule of, or regular bi-weekly payments to be settled at the end of the cost reporting year (that is, a periodic interim payment type approach).

Response: Under § 413.64(k)(3), hospitals have been paid special interim payments for capital costs. With the incorporation of capital costs into the prospective payment system, the hospital-specific portion of capital payments will continue to be paid on a special interim payment basis, as described in § 413.64(k)(3). Payment of the Federal portion of capital payments depends on whether the hospital is entitled to periodic interim payments. (Under section 1815(e) of the Act, as added by section 9311(a) of Pub. L. 99-509, with certain exceptions, payment for inpatient hospital services of hospitals subject to the prospective payment system is no longer made on a periodic interim basis, but rather on the basis of bills actually submitted.) If a hospital meets the exceptions criteria under section 1815(e) of the Act, the hospital may receive periodic interim payments for the Federal portion of its capital payments. Otherwise, the hospital will be paid on the basis of bills actually submitted for the Federal portion of its capital payments, in the same manner as payment is made for other inpatient operating costs.

Comment: Several commenters stated their agreement with our proposal not to adjust standardized capital payment rates for the offset to interest expenses on funded depreciation. However, one commenter suggested that the capital rates should reflect an amount for return on equity capital.

Response: As we pointed out in the June 3, 1986 proposed rule (51 FR 19978), payments for a return on equity capital must be treated in accordance with section 1886(g)(2) of the Act, as amended by section 9107 of Pub. L. 99–272, and are being phased-out separately over a three-year period. We refer the

reader to a June 4, 1987 final rule (52 FR 21216).

Comment: One commenter requested that the final regulations clearly state which capital payment policy would apply to sole community hospitals for cost reporting periods beginning on and after October 1, 1990.

Response: In light of the statutory prohibition excluding sole community hospitals from inclusion in prospective capital payment rulemaking at this time, we expect to announce rules for sole community hospitals at an appropriate time prior to the expiration of that exemption.

#### III. Summary of Final Rule

Under this final rule, hospitals subject to the prospective payment system. other than SCHs, will begin receiving capital payments on a prospective payment basis effective with cost reporting periods beginning on or after October 1, 1987, based on a combination of a Federal capital rate and a hospitalspecific portion during ten-year and seven-year transition periods for plant/ fixed equipment and moveable equipment, respectively. This will continue until capital payments are fully integrated into the prospective payment system with cost reporting periods beginning on or after October 1, 1997. Following is a summary of the methodology we have used in establishing the capital prospective payment rates and of other decisions affecting those payment rates.

#### A. Determination of Federal Capital Payment Rates

Step 1—Split of Plant and Fixed Equipment From Moveable Equipment

Inpatient capital cost data from FY 1984 Medicare cost reports (latest available audited data) were used in the determination of the Federal capital payment rates. In determining the split of plant and fixed equipment from moveable equipment, ratios of plant and fixed equipment costs to the total capital costs, and of moveable equipment costs to the total of capital costs, were developed for each hospital. Capital costs for departments as shown on the Medicare cost report, such as nursery, other reimbursable cost centers (for example, home dialysis), and nonreimbursable cost centers were excluded from these totals because these costs do not represent inpatient capital costs for Medicare payment. Since a split of plant/fixed equipment and moveable equipment could not be determined for directly assigned capital costs at the time the NPRM was

published, directly assigned capital costs were treated as moveable equipment costs only for those hospitals that did not separately report capital costs in the moveable equipment cost center.

For those cases in which capital costs were reported in both the moveable equipment cost center and the directly assigned cost center, the directly assigned costs were not used in the ratio calculations. That is, we assumed that the split of plant and fixed equipment from moveable equipment in the directly assigned cost category is consistent with the split in the plant and fixed equipment from moveable equipment cost centers.

Inpatient capital costs apportioned to the Medicare program were split between plant/fixed equipment and moveable equipment, using the ratios as determined above for each hospital.

Through additional audit efforts, we collected further information on directly assigned capital costs from a sample of hospitals for their FY 1984 cost reports to disaggregate these costs between plant/fixed equipment and moveable equipment, more precisely.

The study results show that the majority of the costs classified to the directly assigned category in the Medicare cost report were for moveable equipment costs. Based on those results, we are adjusting the plant/fixed equipment and moveable equipment ratios determined for each hospital by apportioning directly assigned capital costs in the following manner:

#### [In percent]

	Urban	Rural
Ratio of plant/fixed equipment costs to	ST THE	
total capital costs	28.3	18.2
Ratio of moveable equipment costs to	63363	
total capital costs	71.7	81.8

#### Step 2—Adjustments and Standardizations

#### General

The Federal capital payment rates calculated separately for plant/fixed equipment and for moveable equipment from Federal FY 1984 cost reporting data are adjusted, updated, standardized and computed as separate national averages for urban and rural hospitals, in the 50 States and the District of Columbia. (For background, refer to the NPRM published on May 19, 1987 (52 FR 18846).)

#### Puerto Rico

We developed a separate adjusted average payment rate for urban and

rural hospitals in Puerto Rico using FY 1984 Puerto Rico hospital data and, in general, using the same methodology as that used for the 50 States and the District of Columbia.

#### Standardization

We standardized the Medicare capital plant/fixed equipment and moveable equipment costs of each hospital to eliminate cost variations due to differences in case mix complexity, indirect medical education, and disproportionate share payments (and, for the moveable capital costs of hospitals in Alaska and Hawaii, a cost-of-living adjustment, and for plant/fixed capital costs, a construction cost index adjustment).

We are required to compute urban and rural averages as provided by sections 1886 (d)(2)(D) and (d)(3)(D) of the Act.

#### Local Cost Variations

In recognition of the variations in construction cost among areas, a construction cost index (presented in Appendix B of this document) will be applied in making payments for plant/ fixed equipment. This construction cost index applies only to the Federal payment portion for plant/fixed equipment since such variations are already recognized on an individual hospital basis in determining the hospital-specific portion using a "rolling base." The construction cost index will be applied to standardize the plant/ fixed equipment cost data in the same manner that cost data are standardized for case mix, indirect medical education and disproportionate share. (For a detailed description of the methodology used to compute the construction index, see the May 1987 NPRM (52 FR 18846).)

#### Step 3—Updating

The results from step 2 (the standardized average fixed and moveable costs per case for each hospital) are updated through FY 1987 using the estimated rate of increase in actual inpatient hospital capital costs.

The following update factor percentages were used to establish the capital rates set forth in Appendix A of this document:

Fiscal year	Update factor percent- ages
1985	17.29 10.86
1987	7.06
1988	6.83
1989	5.86

In light of the requirement that aggregate payments for inpatient capital costs under a prospective payment system approximate aggregate payments for inpatient capital costs under cost reimbursement subject to the reductions under section 1886(g)(3) of the Act, we updated hospitals FY 1984 costs per case through FY 1987 by the estimated actual increase in capital costs per case, as shown above, for purposes of establishing the rates. We will also update costs for FY 1988 and FY 1989, as shown above, for purposes of estimating the payments that would be made, subject to the reductions under section 1886(g)(3) of the Act, under reasonable cost principles, as described in further detail in the May 1987 NPRM (52 FR 18849). (The latter estimate will form the basis of the budget neutrality adjustment to the rates.)

For FY 1990 onward, the same update factor methodology will apply to the capital rates as to all other inpatient hospital operating rates.

This methodology will account for-

- Increases in the hospital market basket:
- Plant/fixed equipment separately from moveable equipment;
  - · Cost-effective technologies;
  - · Improved practice patterns:
  - · Pricing of depreciable assets:
  - Consideration of interest expense;
     nd
  - · Productivity.

#### Step 4—Separate Averages

The amounts resulting from step 3 are then used to compute average standardized rates for plant/fixed equipment and moveable equipment, for all urban hospitals and for all rural hospitals in the United States and the District of Columbia, and for urban and rural hospitals in Puerto Rico. The national urban and rural averages are discharge-weighted in the same manner as the prospective payment rates for other inpatient hospital operating costs that will be in effect with discharges occurring on or after October 1, 1987.

#### Step 5—Reducing for Outliers

In accordance with section 1886(d)(5)(A) of the Act, we are amending the current outlier policy (in 42 CFR Part 412, Subpart F) by adding capital to the pool set aside for outliers. The average standardized rates for plant/fixed equipment and moveable equipment resulting from step 4 are reduced by the proportion (estimated by HCFA) of the amount of payments that, based on the total amount of the Federal capital payments for urban hospitals and the total amount of the Federal

capital payments for rural hospitals, are additional payments for outlier cases.

We are providing that payment for capital day outliers (extended length-of-stay cases) will be determined based on the same provisions in effect for noncapital day outliers (§ 412.82). We are providing that payment for capital cost outliers (extraordinarily high-cost cases) will be determined based on both total inpatient operating costs for each case including capital. We are amending § 412.84 to provide that payment for high capital cost cases may occur only when the inpatient operating costs including capital exceed the cost outlier threshold.

The day outlier and cost outlier criteria are revised as follows (note that criteria are the same for capital outliers and noncapital (that is, other inpatient operating cost) outliers):

- For FY 1988, we are setting the day outlier threshold at the lesser of 18 days or two standard deviations, and the cost outlier threshold at the greater of two times the Federal rate for the DRG, or \$14,000.
- We are revising the national ratio of cost to charges used to compute a hospital's cost outlier payments from .66 to .71 to reflect the inclusion of capital costs.

We note that the outlier methodology reflected in this final rule is based on the outlier methodology for other inpatient operating costs as described in the final rule on changes to the inpatient hospital prospective payment system, which is published elsewhere in this issue of the Federal Register.

With the inclusion of capital in the outlier computation, we are providing two examples to illustrate the computation of day outliers and cost outliers. The two examples are applicable to hospitals with cost reporting periods that occur on the same basis as the Federal fiscal year (that is, October 1, 1987) and, therefore, will begin receiving payment for capital under the prospective payment system beginning October 1, 1987.

For hospitals with cost reporting periods that occur after October 1, 1987, capital costs continue to be paid on a pass-through basis as of October 1, 1987 through the end of the hospital's cost reporting period that began in FY 1987. For those hospitals, outlier payments

will continue to be paid for other inpatient operating costs as described in the examples in the September 3, 1986 final rule (51 FR 31523).

The following is an example of how additional payment would be determined for a day outlier (which does not qualify as a cost outlier) in FY 1988.

Hospital X is a small central city teaching hospital located in the San Francisco MSA. Hospital X is entitled to an indirect medical education adjustment of 7.871 percent as well as a disproportionate share adjustment of five percent. Mrs. Smith is admitted to hospital X on October 3, 1987 and is discharged October 31, 1987. Mrs. Smith's stay is classified in DRG 31. Because Mrs. Smith's 28 day stay exceeds the 22 day length-of-stay outlier threshold for DRG 31, hospital X is eligible for payment for six outlier days in addition to the otherwise applicable prospective payment. The amount of Hospital X's outlier payment (excluding the usual Federal payment that applies to both outliers and non-outlier cases) is calculated as follows:

Step 1: Computation of Federal Rate (excludes capital, indirect medical education (IME), and disproportionate share hospital (DSH) payments)

National Urban Standardized	Amounts:
Labor-related	\$2,337.09
Nonlabor-related	828.12
San Francisco Wage Index	
DRG 31 Relative Weight	
.6550 × [2337.09 × 1.4946 + 828	

Step 2: Computation of Federal Capital Payments

A. Plant/fixed Equipment:	
Federal Rate\$180.0	13
San Francisco Construction Cost	
Index1.04	
DRG 31 Relative Weight	0
Federal Portion of Plant/Fixed	
Equipment Capital Rate5	%
$.6550 \times (180.03 \times 1.043) \times .05 = \$6.15$	
B. Movable Equipment:	
Federal Rate\$122.3	9
DRG 31 Relative Weight	0
Federal Portion of Movable Equipment	
Capital Rate5	%
$.6550 \times (122.39) \times .05 = $4.01$	

Standardized cost

C. Capital Outlier Payment Amount: 6.15 + 4.01=\$10.16

Step 3. Payment Amount, Including Capital

DRG 31: 2830.34 + 10.16 = \$2840.50

Step 4. Computation of Day Outlier Payments

DRG 31:
Geometric Length of Stay 4.2 days
Outlier Threshold
Outlier Days28 days length of stay minus 22
day threshold=6 days
Marginal Cost Factor
Outlier Payment (excluding IME & DSH
adjustment) = # of outlier days × (Total
Federal Payment + Geometric length of
stay for DRG) × Marginal cost factor
6 × (2840.50 ÷ 4.2) × .60=\$2437.71
Step 5: Computation of IME and DSH

Step 5: Computation of IME and DSH adjustment for Day Outliers

THE RESERVE THE PROPERTY OF TH	
IME Adjustment Factor	07871
DSH Adjustment Factor	
Outlier Payment	
IME Outlier Adjustment:	
2437.71 × .07871=\$191.87	
DSH Outlier Adjustment:	
2437.71 × .05=\$121.89	

Step 6. Total Day Outlier Payments

Regular	2437.7
ME	191.8
OSH	121.89

The following is an example of how the additional payment would be determined for a high cost outlier in FY 1988. Same facts as in the day outlier example with the exception that Mrs. Smith's length of stay was 16 days and she incurred total billed charges of \$100,000.

Step 1: Computation of Hospital X's Standardized Costs (Includes Capital 2)

Billed Charges—\$100,000 National Ratio of Cost to Charges²—.71 IME Adjustment Factor—.07871 DSH Adjustment Factor—.05

= \$100,000 1+(.07871+.05) ×.71=\$62,903.67

² This factor reflects the inclusion of capital costs and the exclusion of interest income on funded depreciation as described in the June 3, 1986 NPRM (51 FR 20029).

Step 2:	Deter	rmina	tion	of	Cost	Outlier
Thresh	olds					

Computation 1—(Based on Federal R	ate)
DRG 31 Federal rate excluding	
capital	\$2830.34
DRG 31 Federal capital payment ra	

Federal Rate including capital for threshold computation—\$2830.34 + 203.16=\$3033.50

Federal Rate, doubled=2 × \$3033.50=\$6067.00

Computation 2—Based on Adjusted Standard Cost Outlier Threshold

Standard Cost Outlier Threshold—\$16,000 Labor-related share 3—68.632%

San Francisco MSA wage index—1.4946 Nonlabor-related share, excluding capital 3— 23.628%

Nonlabor-related share; capital only3— Plant/Fixed Equipment Share—5.761% Moveable Equipment Share—1.979% Construction Cost Index (San Francisco)—

Adjusted Cost Outlier Threshold including capital =  $(16,000 \times .68632 \times 1.4946) + (16,000 \times .23628) + (16,000 \times .05761 \times 1.043) + (16,000 \times .01979) = $21,470.90$ 

Applicable cost outlier threshold... L\$21,470.90

#### Step 3: Calculation of Cost Outlier Payment

Outlier Cost—\$62,903.67 - 21,470.90 = \$41,432.77

Capital portion of outlier cost
—Plant/Fixed Equipment:

\$41,432.77 × .05761 = \$2386.94 Federal portion of Plant/Fixed Equipment Rate:

Federal Plant/Fixed Equipment Portion of Outlier Cost \$2,386.94 × .05 = \$119.34

#### -Moveable Equipment:

\$41,432.77 × .01979 = \$819.95 Federal Portion of Moveable Equipment Rate: 5%

Federal Movable Equipemnt Portion of Outlier

Cost:  $\$819.95 \times .05 = \$41.00$ Outlier Cost Excluding Capital

\$41,432.77 - (2386.94 + 819.95) = \$38,225.88

Marginal Cost factor.....
Outlier payment—capital and noncapital portions

 $(\$119.34 + 41.00 + 38,225.88) \times .60 = \$23,031.73$ 

Step 4: Cost outlier payment for indirect medical education costs

Percent add-on for indirect medical education—7.871%

Indirect medical education cost outlier payment—\$23,031.73 × .07871 = \$1,812.82

Step 5: Cost outlier payment adjustment for disproportionate share hospital (DSH)

Step 6: Total cost outlier payments

Regular	\$23,031.73
Indirect Medical Education	1,812.82
Disproportionate Share	1,151.69
Total	25,996.24

Step 6—Reducing for additional payments for financially disadvantaged hospitals

Under section 1886(d)(5)(C)(iii) of the Act, we are reducing the average standardized rates for plant/fixed equipment and moveable equipment resulting from step 4 to obtain a five percent pool of funds. This reduction represents the amount necessary to generate two pools of funds to be paid to urban hospitals and rural hospitals that meet the criterion indicating that they are financially disadvantaged under the prospective capital payment system.

We are adding § 412.68 stating that the eligibility criterion is based on whether a hospital's actual, allowable inpatient capital costs not paid through the hospital-specific portion are 175 percent or more than the hospital's total Federal capital payments received based on the Federal prospective capital rates. The amount of the additional capital payment for a hospital meeting the eligibility criterion will be equal to 70 percent of the difference between 175 percent of the hospital's Federal capital payments received based on the Federal prospective capital rates during the transition period and its actual allowable inpatient cost not paid through the hospital-specific portion. We will announce prior to October 1 of each year of the transition period any changes to the threshold level, if needed, to maintain the five percent pool.

#### Step 7—Budget neutrality

Section 1886(g)(3)(C)(ii) of the Act requires that, effective with cost reporting periods beginning during FY 1988, Medicare capital payments under

the prospective payment system shall approximate the amount that would have been made (taking into account the seven percent reduction in capital payments in FY 1988 under section 1886(g)(3)(A)(ii) of the Act) without such inclusion into the prospective payment system (that is, be budget neutral). Since we assumed that the seven percent reduction to capital payments applied with or without the incorporation of capital into the prospective payment system, we did not reduce the amounts resulting from steps 5 and 6 by seven percent. In like manner, since sole community hospitals are excluded from a capital prospective payment system (section 1886(g)(3)(C)(i) of the Act) through cost reporting periods beginning before October 1, 1990, we did not adjust the amounts resulting from steps 5 and 6 for their costs.

We are adjusting the amounts resulting from steps 5 and 6 (by a budget neutrality adjustment factor of 1.0357) so that capital payments during FY 1988 under the capital prospective payment system equal the capital payments that would have been paid for the same time period in the absence of this system (that is, under the reasonable cost reimbursement system). Taking into consideration the different blends of the hospital-specific and Federal portions applied to plant/fixed equipment and moveable equipment in FY 1988, the budget neutrality equation is as follows:

.95 HSP Fixed + .05 Fed Fixed + .95 HSP Mov + .05 Fed Mov = CRC Fixed + CRC Mov

## where-

 HSP Fixed and HSP Mov represent the total hospital-specific payments made for plant/fixed equipment and moveable equipment during FY 1988 under the capital prospective payment system;

 Fed Fixed and Fed Mov are the total Federal payments for plant/fixed equipment and moveable equipment during FY 1988 under the capital prospective payment system; and

 CRC Fixed and CRC Mov are the capital costs for plant/fixed equipment and moveable equipment that would have been paid under the reasonable cost reimbursement system.

Since the hospital-specific capital payments are based on actual costs in FY 1988, the budget neutrality equation can be modified as follows:

.05 Fed Fixed + .05 Fed Mov = .05 HSP Fixed + .05 HSP Mov The terms on both sides of the equation were estimated for each hospital applying the payment rules

described elsewhere in this final rule.

³ These market basket weights reflect updated 1986 market basket components, including capital, which are based on 1982 cost data as described in the June 3, 1986 proposed rule (51 FR 19985–19988).

For each hospital, the hospital-specific payments were adjusted to the midpoint of the hospital's FY 1988 fiscal year. The Federal payments were estimated using rates adjusted to the midpoint of FY 1987. In both cases, these adjustments were based on estimates of the actual rate of increase in capital costs per admission derived from AHA data. As a result of the one year difference between the updating of the hospital-specific and Federal variables, the adjustment factor defined below accounts both for capital cost differences between 1987 and 1988 and for the budget neutrality requirement.

Federal payments were computed using case-mix indexes for FY 1986.
Since all estimates were made on a per discharge basis, the number of discharges has no effect on the value of the adjustment factor.

To calculate the required adjustment to the Federal rates, we substituted the adjusted Federal payments for the Federal terms in the equation above. Let the adjustment factor applied to the Federal rates be k. Then the adjusted Federal payments are k*Fed fixed and k*Fed Mov. Solving the resulting equation for k yields the following:

k=(.05 HSP Fixed + .05 HSP Mov) / (.05 Fed Fixed + .05 Fed Mov)

#### B. Determination of the Hospital-Specific Portion of the Capital Payment

The hospital-specific portion of the capital payment during the transition period will be made by determining Medicare's share of the actual allowable capital costs of hospital plant/fixed equipment and of moveable equipment apportioned to inpatient areas for each transition year as determined under Medicare's principles of reimbursement that implement section 1861(v) of the Act and Subpart G of 42 CFR Part 413. We then determine the hospital-specific amount by reducing those costs by the percentage factors (seven and ten percent) mandated for FYs 1988 and 1989, respectively, and the appropriate blend percentages.

## Step 1—Allocation of equipment

The amount of Medicare allowable capital costs of plant/fixed equipment and of moveable equipment, separately apportioned to inpatient hospital services, will be determined pursuant to \$\$ 412.113(a) and 413.130. Due to the importance of splitting fixed/plant from moveable equipment, we are providing a detailed list for classifying the current and future items of hospital capital in Appendix C. The HCFA list is an adaptation of the list of the American Hospital Association (AHA) 1987

Edition and is an extensive, but not exhaustive list of items of plant/fixed equipment, and of moveable equipment. (We noted in the May 1987 NPRM that we have added extracorporeal shockwave lithotripters and magnetic resonance imaging equipment to the AHA list of equipment under the moveable equipment category.) We will update the list periodically as necessary through notices published in the Federal Register. We also note that the definitions contained in the Provider Reimbursement Manual (HCFA Pub. 15-1, Chapters 1 and 2) will be used to classify equipment not specified on the list. We will also allow a hospital that has been classifying items differently than the classifications under the list in Appendix C of this final rule, or as defined in § 412.67(d), to continue its current classification practice if the present classification was in effect prior to its cost reporting period beginning on or after October 1, 1987, and the hospital demonstrates to the fiscal intermediary that the classification of equipment in question is based on longstanding past practices and is not for the purpose of affecting prospective capital payments.

#### Step 2—Apportionment

The Medicare allowable capital costs of plant/fixed equipment and of moveable equipment, apportioned to inpatient hospital services, are reduced by the appropriate percentage factor pursuant to section 1886(g)(3) of the Act, as added by section 9303 of Pub. L. 99–509. The capital costs thus reduced by seven percent and ten percent in FYs 1988 and 1989, respectively, are then multiplied by the appropriate hospital-specific blend percentage applicable to the pertinent transition cost reporting period.

#### C. Additional Payments

We are directed by section 1886(d)(5) of the Act to provide additional payments for outliers (section 1886(d)(5)(A)), indirect medical education (section 1886(d)(5)(B)), and disproportionate share adjustments (section 1886(d)(5)(C)) for hospitals that are under the prospective payment system. As a result, we are required to add to the current inpatient operating payment the portion of the Federal Capital payment for those additional payments that we are including in the prospective payment rate. In addition, pursuant to the Secretary's discretionary exceptions and adjustments authority under section 1886(d)(5)(C)(iii) of the Act, we will be providing additional payments to hospitals that are financially disadvantaged by the

changeover to the prospective capital payment system.

#### 1. Payments for Outliers

The pool of funds for day and cost outlier payments is increased by reducing the average standardization Federal capital payment rates by the same factors used to reduce inpatient operating prospective payment rates. In determining the amount of a day outlier payment, we are using the same methodology currently used for inpatient operating costs. That is, the Federal capital and inpatient operating standardized amounts, adjusted to reflect the construction cost index and the wage index (as appropriate), and also to reflect the appropriate blends, are multiplied by the applicable DRG weighting factor. The resulting adjusted standardized amounts for capital and other inpatient operating are added together, and the result divided by the geometric average length-of-stay figure for the DRG. This amount is then multiplied by the marginal cost factor, and the resulting amount will be paid to the hospital for each day of care beyond the day outlier threshold. We refer the reader to section III.A. step 5 of the preamble for a further discussion and examples of the outlier provisions.

In determining the amount for a cost outlier payment, we are using the same methodology currently used for other inpatient operating costs. Since we are including capital costs, as part of total inpatient operating costs, we will make any appropriate changes to the cost outlier thresholds to reflect additional capital costs. The plant/fixed capital portion of the thresholds will be adjusted by the construction cost index in the same manner as the other inpatient operating portion is adjusted by the hospital wage index. In addition, the national cost-to-charge ratio used in determining cost outlier payments will be increased to reflect capital costs.

We reiterate that capital outlier payments will not be paid until a hospital is subject to prospective payments for capital, that is, effective for discharges in cost reporting periods beginning on or after October 1, 1987. For example, if a hospital's cost reporting period in FY 1988 begins January 1, 1988, capital outlier payments would be made, if applicable, for discharges occurring on or after January 1, 1988. Therefore, adjustments to outlier payments for capital would not be available to that hospital until January 1, 1988. In addition, changes to both cost and day outliers criteria and capital outlier payments would apply on a Federal fiscal year basis in order to

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maintain consistency with the outlier policy for other inpatient operating costs under the prospective payment system. This means that changes in outlier criteria and payments announced next year to be effective for Federal FY 1989 will apply to the January 1, 1988 hospital as of October 1, 1988.

In summary, we are making appropriate changes to §§ 412.82 and 412.84 to incorporate capital into the outlier criteria and payments.

#### 2. Payments for Indirect Medical Education

We adjusted the average standardized amounts to account for indirect medical education payments. Hospitals that are eligible for indirect medical education payments under § 412.118 will receive an additional payment to their capital payments for indirect medical education costs to be calculated in the same

manner as other inpatient operating indirect medical education payments are determined

3. Payments for Hospitals that Serve a Disproportionate Share of Low-Income Patients

We adjusted the average standardized amounts to account for the costs of hospitals that serve a disproportionate share of low-income patients. Hospitals that meet the disproportionate share criteria under § 412.106 will receive an additional payment to their capital payments for disproportionate share costs to be calculated in the same manner as other inpatient operating disproportionate share payments are determined.

4. Payment Adjustment for Hospitals that Meet the Eligibility Requirements for an Exceptions Process

We adjusted the average standardized amounts to account for the cost of increasing the Federal capital payment amount for hospitals that meet the eligibility criteria established for hospitals that are financially disadvantaged during the transition period by the level of their capital payments in comparison to their actual inpatient hospital capital cost per discharge. As discussed above, we are providing a reserve pool of funds necessary to make these payments.

## D. Total Capital Payment

The total capital payment to a hospital for any particular cost reporting period will be a total of the following components:

· The total of the hospital-specific amounts for discharges occurring in a period according to the following formula:

Medicare's Share of Aliowable Cost for X Applicable Percentage Reduction X Hospital-Specific Portion Transition Plant and Fixed Equipment. Factor.4. Blend Percentage. -Plus-

Medicare's Share of Allowable Cost for X Applicable Percentage Reduction X Hospital-Specific Portion Transition Moveable Equipment. Factor.4. Blend Percentage.

 The aggregate Federal portion payment amounts for plant/fixed

Federal Rate Per

Discharge.5.

equipment and for moveable equipment, for discharges occurring during the

pertinent cost reporting period based on the following formula:

Total of Plant and Fixed Equipment Federal Rate Per Discharge.5.

X Area Construction Index.

X DRG Weight ..... X Any Applicable Adjustment.6.

X Federal Transition Blend Percentage.

-Plus-Moveable Equipment X DRG Weight ...... X Any Applicable

X Federal Transition Blend Percentage.

-Plus-

Any Applicable Outlier and Exceptions Payments.

#### E. New Hospitals

As a result of using a "rolling" base to determine the hospital-specific portion of the capital payment during the transition period, no special provision will be made for new hospitals that become subject to the prospective payment system upon entering the Medicare program. A new hospital is subject to the capital payment transition blends in effect at the beginning of its cost reporting period when it enters the program. Thus, a hospital entering the program (that is, newly participating in the Medicare program, under present or previous ownership) on January 1, 1993

will be paid at 70 percent of its hospitalspecific costs and 30 percent of its Federal rate for plant/fixed equipment and 100 percent of its Federal rate for moveable equipment.

#### F. Sole Community Hospitals

As prescribed in section 1886(g)(3)(C)(i) of the Act, sole community hospitals will continue to be paid under the reasonable cost methodology described in section 1861(v)(1) of the Act with respect to capital costs for cost reporting periods beginning before October 1, 1990.

## G. Capital Expenditure Agreements

Section 1886(g)(1) of the Act provides that, if legislation concerning payment for capital costs for inpatient hospital services is not enacted before October 1, 1987, no payment may be made for capital costs of capital expenditures (as defined in section 1122(g) and except as provided in section 1122(j) of the Act) for inpatient hospital services in a State (including the District of Columbia and Puerto Rico as defined in sections 210(h) and 1861(x) of the Act) if such expenditures are obligated after September 30, 1987, unless the State or jurisdiction has an agreement with the

^{*} Represents the reduction applicable for FYs 1988 and 1989 as provided under section 1886(g)(3) of the Act.

hincludes the reductions and budget neutrality adjustments applicable for FYs 1988 and 1989 as provided under section 1886(g)[3] of the Act. For example, indirect medical education, disproportionate share.

Secretary under section 1122(b) of the Act and under such agreement the State recommended approval of the capital

expenditure.

The deadline in section 1886(g)(1) of the Act for legislation to avoid invocation of the section 1122 requirements was extended to October 1, 1987, by section 206 of Pub. L. 99-349, enacted July 2, 1986. Subsequently, under section 9303 of Pub. L. 99-509, Congress enacted section 1886(g)(3) of the Act, which requires certain reductions in the payments for capital costs for inpatient hospital services. We stated in the NPRM that we believe that section 1886(g)(3) constitutes the necessary legislation required under section 1886(g)(1) "respecting the payment . . , for capital-related costs for inpatient hospital services," thus nullifying the requirement of section 1886(g)(1).

#### H. Interest Expense

As we stated in the May 1987 NPRM, interest expense, as described in § 413.153, is an integral part of capital costs. Under the current Medicare capital payment system (reasonable costs), it is important for hospitals to distinguish operating interest from capital interest appropriately since interest on funds borrowed for operating expenditures is included under the inpatient hospital prospective payment system, and therefore is not a passthrough, while interest on funds borrowed for capital expenditures is paid for on a reasonable cost passthrough basis. We proposed in the NPRM to continue the application of this principle without change throughout the transition period. We will apply this principle and Medicare's principles of reimbursement to establish the amount and appropriate split of capital interest expense for purposes of determining the hospital-specific portion of each hospital's prospective capital payment during the transition period, in the manner described below.

# 1. Treatment of interest expense

We are determining interest expense allocation based on current regulations (§ 413.153) and program guidelines (see section 3202.1 of the Provider Reimbursement Manual (HCFA Pub. 15–1)). To be allowable as a Medicare expense, interest must be—

 Supported by evidence of an agreement that funds were borrowed and that payment of interest and repayment of funds are required;

Identifiable in the hospital's records;

 Related to the reporting period in which the costs are incurred; and

 Necessary and proper for the operation, maintenance, or acquisition of the hospital's facilities.

To support the existence of a loan, the hospital must have available a signed copy of the loan contract, which should contain the pertinent terms of the loan. If the lender does not customarily furnish a copy of the loan contract, correspondence from the lender stating the pertinent terms of the loan would be acceptable. If interest expense has been determined to be allowable and the interest expense records are maintained physically away from the hospital's premises, such as a county treasurer's office, these records will be deemed to be those of the hospital. This is applicable when bond issues have been specifically designated for the construction or acquisition of hospital facilities and the financial records relative to the bond issue are maintained by some governmental body.

Once the allowable interest expense on capital indebtedness is determined, the interest expense is to be classified to plant/fixed equipment or to moveable equipment using the current Medicare principles of reimbursement for classifying interest.

If a loan is obtained to finance the purchase of a facility or moveable equipment, the interest expense is classified to plant/fixed equipment or to moveable equipment, as appropriate.

If a loan is obtained to finance the purchase of facilities and various equipment items, the interest expense must be distributed among the assets the loan covers based on the purchase price of the acquisitions.

#### Example:

Assets purchased	Purchase cost
Buildings & Fixtures Moveable Equipment	\$240,000 60,000
Total	300,000

Of the \$300,000 purchase price, assume the hospital borrowed \$270,000 for buildings, fixtures, capital improvements and moveable equipment at 10 percent annual interest. Thus, annual interest on the loan is equal to \$27,000. The allocation to plant/fixed equipment and to moveable equipment is shown below:

Building Fixtures  $\frac{\$240,000}{\$300,000} \times 270,000 \times 10\% = \$21,600$  Moveable Equipment  $\frac{\$60,000}{\$300,000} \times \$270,000 \times 10\% = \underbrace{5,400}_{\$27,000}$  Total Interest Expense

If a loan is obtained to finance the purchase of a facility and equipment and the loan exceeds the asset value of the acquisitions, the interest expense on that portion of the loan in excess of the asset value of the acquisitions is not considered capital. The portion of the interest expense on the asset value of the acquisitions must be distributed among the assets of the loan as described in the example above.

There are some cases in which a hospital may, for a variety of reasons, undertake advance refunding (that is, to replace existing debt prior to its scheduled maturity with new debt). The revenues and expenses associated with the advance refunding are treated in accordance with the principles set forth in section 233 of the Provider Reimbursement Manual. The allocation of interest expense on the new debt will

be dependent upon the allocation used under the old debt.

If a hospital has consolidated various individual debts through advance refunding, the interest expense on the new debt will be allocated to the appropriate accounts based on the old debt balances that were refinanced.

Further refinements and clarifications to the cost-finding rules and the cost-reporting methodology will be developed through HCFA's administrative issuances system.

# 2. Treatment of Interest Income from Funded Depreciation

To minimize the disruption in a hospital's cash flow during the initial part of the transition period, we will not offset the interest expense for income earned on funded depreciation.

#### I. Revaluation of Assets

Under section 1861(v)(1) of the Act, for hospital acquisitions which involve revaluation of assets, the new depreciation value of the purchased asset is limited to the lesser of the purchase price or original book value of the asset. When the sale price of the asset exceeds the net book value, Medicare recaptures its proportion of previous depreciation payments from the seller.

Under this final rule, fixed and moveable capital expenses will be phased-in using different transition schedules. For fixed assets, the amount of the capital expenses after adjustments for gains and losses due to changes of ownership will be equal to the percentage of fixed assets (as provided in the proposed transition period) that are subject to the hospitalspecific portion of capital costs. Similarly, for moveable assets, the capital expenses after adjustment for gains and losses due to changes of ownership will be the percentage of moveable equipment (as provided in the proposed transition period) subject to the hospital-specific capital payment.

#### J. Hospitals and Units Not Subject to the Prospective Payment System

Capital costs for hospitals and units excluded from the prospective payment system will continue to be paid on a reasonable cost basis.

#### IV. Other Required Information

#### A. Impact Statement

See Appendix D for the Regulatory Impact Analysis.

#### B. Paperwork Reduction Act

In the May 1987 NPRM we stated that § 412.65(b) of this rule contains information collection requirements subject to approval by the Executive Office of Management and Budget under section 3507 of the Paperwork Reduction Act (44 U.S.C. 3507). The requirements of this final rule do not impose any information collection requirements that must be approved under the Paperwork Reduction Act. Therefore, the final rule need not be reviewed by EOMB for that purpose.

#### List of Subjects in 42 CFR Part 412

Health facilities, Medicare. 42 CFR Part 412 is amended as

#### PART 412—PROSPECTIVE PAYMENT SYSTEM FOR INPATIENT HOSPITAL SERVICES

A. The authority citation for Part 412 continues to read as follows:

Authority: Secs. 1102, 1122, 1871, and 1886 of the Social Security Act, as amended (42 U.S.C. 1302, 1320a-1, 1395hh, and 1395ww).

B. The Table of Contents of Part 412 is amended by adding the titles of new §§ 412.65 through 412.68 to Subpart D, and by adding § 412.214 to Subpart K to read as follows:

#### Subpart D—Basic Methodology for Determining Federal Prospective Payment Rates

Sec. * * * * *

412.65 Incorporation of capital payments into the prospective payment system.
 412.66 Federal capital rates beginning

during and after Federal fiscal year 1988. 412.67 Phase-in period and methodology for capital payments.

412.68 Additional payments for capital costs.

# Subpart K—Prospective Payment System for Hospitals Located in Puerto Rico

Sec. 412.214 Capital payments. * * * * *

C. Subpart A is amended as follows:

#### Subpart A-General Provisions

 Section 412.1 is amended by revising paragraph (a) to read as follows:

#### § 412.1 Scope of part.

(a) Purpose. This part implements section 1886(d) of the Act by establishing a prospective payment system for inpatient hospital services furnished to Medicare beneficiaries in cost reporting periods beginning on or after October 1, 1983. Under the prospective payment system, payment for the operating costs of inpatient hospital services furnished by hospitals subject to the system (generally, shortterm, acute-care hospitals) is made on the basis of prospectively determined rates and applied on a per discharge basis. Payment for other costs related to inpatient hospital services (capital costs for cost reporting periods beginning on or after October 1, 1983 and before October 1, 1987, kidney acquisition costs incurred by hospitals with approved renal transplantation centers, direct costs of medical education, and, for cost reporting periods beginning on or after October 1, 1984 and before October 1, 1987, the costs of qualified nonphysician anesthetists' services) is made on a reasonable cost basis. Additional payments are made for outlier cases, bad debts, and indirect medical education costs. Under the prospective payment system, a hospital may keep

the difference between its prospective payment rate and its operating costs incurred in furnishing inpatient services, and is at risk for operating costs that exceed its payment rate.

2. In § 412.2, the introductory language of paragraphs (c) and (d) is republished; a new paragraph (c)(5) is added; and paragraph (d)(1) is revised to read as follows:

## § 412.2 Basis of payment.

. .

* *

(c) Inpatient operating costs. The prospective payment system provides a payment amount for inpatient operating costs, including—

(5) For cost reporting periods beginning on or after October 1, 1987, capital costs as described in Subpart D of this part.

(d) Excluded costs. The following inpatient hospital costs are excluded from the prospective payment amounts and paid for on a reasonable cost basis:

(1) For cost reporting periods beginning on or after October 1, 1983 and before October 1, 1987, capital costs as described in § 413.130 of this chapter; and an allowance for return on equity, as described in § 413.157 of this chapter.

D. Subpart D is amended as follows:

#### Subpart D—Basic Methodology for Determining Federal Prospective Payment Rates

 Section 412.63 is amended by revising paragraph (a)(1) to read as follows:

# § 412.63 Federal rates for fiscal years after Federal fiscal year 1984.

(a) General rule. (1) HCFA determines a national adjusted prospective payment rate for each inpatient hospital discharge in a Federal fiscal year after fiscal year 1984 (including an additional payment, effective with cost reporting periods beginning on or after October 1. 1987, for the incorporation of capital payments as described in § 412.65) involving inpatient hospital services of a hospital in the United States subject to the prospective payment system, and determines a regional adjusted prospective payment rate for such discharges in each region, for which payment may be made under Medicare Part A.

2. New §§ 412.65 through 412.68 are added to read as follows:

### § 412.65 Incorporation of capital payments into the prospective payment system.

- (a) General rule. As described in §§ 412.66 and 412.67, effective with cost reporting periods beginning on or after October 1, 1987, HCFA pays an amount for capital costs for each inpatient hospital discharge, in addition to the Federal rates as determined under § 412.63. In certain cases, HCFA makes additional capital payments as provided in § 412.68.
- (b) Cost reporting periods beginning on or after October 1, 1987 through September 30, 1987. For cost reporting periods beginning during the period October 1, 1987 through September 30, 1997, the capital payment amount is based on a combination of a hospital specific capital portion and a Federal capital rate as determined under §§ 412.66 and 412.67.
- (c) Cost reporting periods beginning after the transition periods end. For cost reporting periods beginning on or after October 1, 1994, the capital payment amount for movable equipment is based solely on a Federal capital rate as determined under § 412.66(h). For cost reporting periods beginning on or after October 1, 1997, the capital payment amount for plant and fixed equipment is based solely on a Federal capital payment as determined under § 412.66(h).

# § 412.66 Federal capital rates beginning during and after Federal fiscal year 1988.

(a) Determining allowable base-year capital costs. HCFA determines the Federal capital rate on the basis of hospitals' Medicare capital costs per discharge, as described in § 413.130 of this chapter, using hospital cost reports from fiscal year 1984 for hospitals in the fifty States and the District of Columbia subject to the prospective payment system, and from short-term acute care hospitals in Puerto Rico.

(b) Separating moveable equipment from plant and fixed equipment. For purposes of the phase-in period, as described in § 412.67, HCFA separates portions of each hospital's inpatient capital costs determined under paragraph (a) of this section that are attributable to moveable equipment from those portions attributable to plant/fixed equipment.

(c) Standardizing the amounts. (1)
HCFA standardizes each portion of
plant/fixed equipment and of moveable
equipment, determined under paragraph
(b) of this section, for each hospital as
follows:

(i) For both moveable equipment and plant/fixed equipment, by—

 (A) Adjusting for resource intensity in case mix among hospitals;

(B) Excluding an estimated amount of indirect medical education payments;

(C) Excluding an estimated amount of the payments for hospitals that serve a disproportionate share of low-income patients; and

(D) Adjusting for the reductions to capital payments under § 413.64(k)(6) of

this chapter.

(ii) For moveable equipment, by adjusting for the effects of higher cost of living payments to hospitals located in Alaska and Hawaii.

(iii) For plant/fixed equipment, by adjusting for the effects of a capital

construction cost index.

(2) Based on the standardizations calculated in paragraphs (c)(1)(i) through (c)(1)(iii) of this section, HCFA determines standardized rates for plant/fixed equipment and for moveable equipment, for hospitals in the fifty States and the District of Columbia subject to the prospective payment system, and for hospitals in Puerto Rico subject to the prospective payment system.

(d) Updating the capital costs. HCFA updates each hospital's adjusted plant/fixed and moveable costs per case determined under paragraph (c) of this

section by-

(1) Updating from fiscal year 1984 through fiscal year 1987 using estimated increases in actual capital costs per case:

(2) Updating for fiscal years 1988 and 1989 using the respective annual estimated increase in actual capital costs per case, as adjusted in accordance with § 413.64(k)(6); and

(3) Projecting for fiscal year 1990 onward the applicable percentage

change under § 412.63(e).

(e) Computing urban and rural averages. HCFA computes a discharge-weighted average of the standardized amounts determined under paragraph (d) of this section for all urban hospitals and for all rural hospitals, as defined in § 412.62(f), in the fifty States and the District of Columbia, and for urban hospitals and rural hospitals in Puerto Rico. HCFA also computes a discharge-weighted average of the urban capital payment rate and the rural capital payment rate for hospitals in Puerto Rico.

(f) Reducing for value of outlier payments and additional capital payments. Based on the total amount of the Federal capital payments for urban hospitals and the total amounts of the Federal capital payments for rural hospitals, HCFA reduces each of the average standardized amounts

determined under paragraph (e) of this section by-

(1) The proportion (estimated by HCFA) of the amount of payments that are set aside for additional payments for outlier cases, as provided under Subpart F of this part; and

(2) Five percent (estimated by HCFA) of the total amount of payments, which is set aside for additional payments for capital costs, as provided under § 412.68.

(g) Application of blending percentages during the phase-in period. For cost reporting periods beginning during the period October 1, 1987 through September 30, 1997, the amounts for plant/fixed equipment and for moveable equipment determined separately under paragraphs (b) through (f) of this section are multiplied by the appropriate phase-in period percentages, respectively, as described in § 412.67(b).

(h) Federal capital payment. (1) Except for sole community hospitals as described in paragraph (h)(2) of this section, the Federal capital payment

equals the product of-

(i) The national capital rates as determined under paragraphs (a) through (g) of this section and § 412.67(b) including an adjustment to the plant/fixed equipment standardized amounts for the construction cost index and an adjustment to the moveable equipment standardized amounts for the higher cost of living for hospitals located in Alaska and Hawaii; and

(ii) The DRG weighting factor determined under § 412.60(b) for each

discharge.

(2) For cost reporting periods beginning before October 1, 1990, sole community hospitals are paid on a reasonable cost basis, as provided under Part 413 of this chapter, for their capital costs.

(i) Additional capital payments. HCFA makes additional capital

payments to hospitals-

(1) That serve a disproportionate share of low-income patients, as described in § 412.106; and

(2) For indirect medical education costs, as described in § 412.118.

# § 412.67 Phase-in period and methodology for capital payments.

(a) Phase-in period. Except for new hospitals and sole community hospitals as described in paragraphs (e) and (f) of this section, respectively, inclusion of payments for capital for plant/fixed equipment and for moveable equipment in the prospective payment rates is to be phased-in over a ten-year period and a seven-year period, respectively, as described in paragraph (b) of this

section. During this period, the capital payment amount is based on a combination of a hospital-specific capital portion and a Federal capital rate as determined in § 412.66. At the end of the transition periods (that is, for discharges occurring in a cost reporting period beginning on or after October 1, 1994, for moveable equipment, and on or after October 1, 1997, for plant/fixed equipment), payment amounts are based entirely on a Federal capital rate.

(b) Blended percentages for capital rates. The blends of the hospital-specific capital portions and the Federal capital rates, for plant/fixed equipment and for movable equipment, are described in the

following tables:

TABLE.—HOSPITAL-SPECIFIC AND FEDERAL RATE PERCENTAGES FOR DETERMINING PHASE-IN PERIOD CAPITAL RATES FOR PLANT AND FIXED EQUIPMENT

Cost reporting period beginning on or after	Hospi- tal- specific capital per- centage	Federal capital per- centage
Oct. 1, 1987	95	5
Oct. 1, 1988	90	10
Oct 1, 1989	85	15
Oct. 1, 1990	80	20
Oct. 1, 1991	75	25
Oct. 1, 1992	70	30
Oct. 1, 1993	60	40
Oct 1, 1994	50	50
Oct. 1, 1995	35	65
Oct. 1, 1996		80
Oct 1, 1997		100

TABLE.—HOSPITAL-SPECIFIC AND FEDERAL RATE PERCENTAGES FOR DETERMINING PHASE-IN PERIOD CAPITAL RATES FOR MOVEABLE EQUIPMENT

Cost reporting period beginning on or after	Hospo- tal- specific capital per- centage	Federal capital per- centage
Oct. 1, 1987	95	5
Oct. 1, 1988	90	10
Oct. 1, 1989	85	15
Oct. 1, 1990	80	20
Oct. 1, 1991	70	30
Oct. 1, 1992	50	50
Oct. 1, 1993	25	75
Oct 1, 1994		100

(c) Hospital-specific capital portion. The hospital-specific capital portion is the hospital's actual allowable Medicare inpatient hospital costs attributed to plant/fixed equipment and to moveable equipment, determined separately, for the applicable phase-in year, multiplied by the appropriate phase-in period percentages described in paragraph (b) of this section, and adjusted for fiscal years 1988 and 1989 by the reductions described under § 413.64(k).

(d) Classification of capital assets as plant/fixed or moveable equipment—(1) General rule. Except as specified in

paragraph (d)(4) of this section, a hospital must classify its capital assets as plant/fixed equipment or as moveable equipment as described in this paragraph for purposes of receiving payment for capital expenditures under the prospective payment system.

(2) Procedures for classifying assets.
(i) HCFA establishes a list under which HCFA assigns capital assets to a plant/fixed equipment category or a moveable equipment category and updates the list periodically as necessary in notices published in the Federal Register.

(ii) Capital assets not specified on the list are assigned to the plant/fixed or moveable equipment categories under the definitions provided in paragraph

(d)(3) of this section.

(3) Definitions. The following definitions apply for purposes of classifying assets not specified on the HCFA list.

(i) "Plant/fixed equipment" means-

(A) A building, which includes, in a restrictive sense, the basic structure or shell and additions thereto (with the remainder being identified as building

equipment); and

- (B) Building equipment, the general characteristics of which are that it is affixed to the building, and not subject to transfer; and that it has a fairly long useful life, but one that is shorter than the useful life of the building to which it is affixed.
- (ii) "Moveable equipment" means equipment that has the following general characteristics:
- (A) A relatively fixed location in the building;
- (B) Capability of being moved as distinguished from building equipment;
- (C) A unit cost sufficient to justify ledger control;
- (D) Sufficient size and identity to make control feasible by means of identification tags; and

(E) A minimum useful life of approximately three years.

(4) Exception for past classification practices. A hospital may request review of the classification of a capital asset that is classified under paragraph (d)(2) or (d)(3) of this section, if the hospital demonstrates to its fiscal intermediary that its classification of the asset during its cost reporting period beginning before October 1, 1987—

(i) Is based on longstanding past practices; and

(ii) Is not for the purpose of affecting Federal capital prospective payments.

(e) Payment rate for newlyparticipating hospitals. (1) If a hospital meets the criteria in paragraph (e)(2) or (e)(3) of this section, it is paid on the basis of the Federal capital portion, as determined in § 412.66, and the hospitalspecific capital portion, as determined in paragraph (c) of this section, using the blending percentages applicable for the Federal fiscal year in which it initially participates in the Medicare program.

(2) The hospital-

- (i) Is newly participating in the Medicare program (under previous and present ownership); and
- (ii) Does not have a 12-month cost reporting period ending on or before September 30, 1987.
- (3) The hospital is under new ownership and documents to the satisfaction of its intermediary that the ownership and occupancy rate requirements described in § 412.74(a)(2) are met.
- (f) Payment rate for sole community hospitals. For cost reporting periods beginning before October 1, 1990, a hospital that meets the criteria in § 412.92(a) for classification as a sole community hospital is paid on a reasonable cost basis, as provided under Part 413 of this chapter, for its capital costs.

# § 412.68 Additional payments for capital costs.

- (a) Threshold for additional payments. During the phase-in periods for capital payments, as described in § 412.67, a hospital may be paid an additional capital payment for plant/fixed equipment or moveable equipment, or both, if the portions of the hospital's actual allowable inpatient capital cost not paid through the hospital-specific portions (as determined under § 412.67) equal or exceed a percent (specified by HCFA in the Federal Register notices published under § 412.8) of its total Federal capital payments received under § 412.66.
- (b) Payment amount. A hospital that meets the criterion described in paragraph (a) of this section may be paid an additional capital payment equal to 70 percent of the difference between—
- (1) The portion of the hospital's total actual allowable inpatient capital cost not paid through the hospital-specific portion as determined under § 412.67(c); and
- (2) The percent (specified by HCFA) of the hospital's total Federal capital payments received under § 412.66.
  - E. Subpart F is amended as follows:

# Subpart F-Payment for Outlier Cases

1. In § 412.82, paragraph (c) is revised to read as follows:

# § 412.82 Payment for extended length-ofstay cases (day outliers).

(c) The per diem payment made under paragraph (a) of this section is derived by first taking 60 percent of the average per diem payment for the applicable DRG, as calculated by dividing the Federal prospective payment rates (noncapital, and effective with cost reporting periods beginning on or after October 1, 1987, capital) determined under Subpart D of this part by the geometric mean length-of-stay for that DRG. The resulting amounts are then multiplied by the applicable Federal portions (capital and noncapital) of the blend as follows:

### FEDERAL NONCAPITAL PORTIONS

Cost reporting periods beginning on or atter	Fed- eral por- tion (per- cent)
October 1, 1983	25 50
October 1, 1985	50
The remaining five months of the cost reporting period	55 75 100

# FEDERAL CAPITAL PORTIONS

Cost reporting periods beginning on or after	Plant and fixed equip- ment (per- cent)	Mov- able equip- ment (per- cent)
Oct 1, 1987	5	5
Oct. 1, 1988	10	10
Oct. 1, 1989		15
Oct. 1, 1990		20
Oct. 1, 1991	25	30
Oct. 1, 1992		50
Oct. 1, 1993	40	75
Oct. 1, 1994		100
Oct. 1, 1995		100
Oct. 1, 1996	80	100
Oct. 1, 1997	100	100

2. In § 412.84, paragraphs (g) and (i) are revised to read as follows:

### § 412.84 Payment for extraordinarily highcost cases (cost outliers).

.

. .

(g) The intermediary bases the cost of the discharge on a certain percentage of the billed charges for covered impatient services. The cost is adjusted further to exclude an estimate of indirect medical education costs, and payments for hospitals that service a disproportionate share of low-income patients, and to include the reasonable charges for nonphysician services billed by an outside supplier in accordance with § 489.23(c)(3) of this chapter.

(1) For discharges in cost reporting periods beginning before October 1, 1987, the factor is 66 percent.

(2) For discharges in cost reporting periods beginning on or after October 1, 1987, the factor is 71 percent.

(i) The additional payment amount is derived by first taking 60 percent of the difference between the hospital's adjusted cost for the discharge (as determined under paragraph (g) of this section) and the threshold criteria established under § 412.80(a)(1)(ii). The resulting amounts are then multiplied by the applicable Federal portions (noncapital, and effective with cost reporting periods beginning on or after October 1, 1987, capital) of the blend as indicated in § 412.82(c).

F. Subpart G is amended as follows:

# Subpart G—Special Treatment of Certain Facilities

1. In § 412.92, paragraph (d) is revised to read as follows:

# § 412.92 Special treatment: Sole community hospitals.

(d) Determining prospective payment rates for sole community hospitals. For all cost reporting periods beginning on or after October 1, 1983, the prospective payment rates for sole community hospitals equal the sum total of the following payment rates:

(1) 75 percent of the hospital-specific base payment rate as determined under \$ 412.73.

§ 412.73;

(2) 25 percent of the appropriate regional prospective payment rate as determined under Subpart D of this part; and

(3) The capital payment as determined under § 412.67(f).

2. In § 412.96, paragraphs (d) and (e) are revised to read as follows:

# § 412.96 Special treatment: Referral centers.

(d) Payment to rural referral centers with 500 or more beds. A hospital that meets the criteria of § 412.96(b)(1) is paid prospective payments per discharge based on the applicable urban payment rates as determined in accordance with § 412.62(j) or § 412.63(f), as adjusted by the hospital's area wage index, and § 412.66(h), as adjusted by the capital construction cost index adopted by HCFA and applicable to the hospital.

(e) Payment to all other rural referral centers. For cost reporting periods beginning on or after October 1, 1984, a hospital that is located in a rural area and meets the criteria of § 412.96 (b)(2) or (c) is paid prospective payments per discharge based on the applicable urban payment rates determined in accordance with §412.62(j) or § 412.63(f), as adjusted by the hospital's area wage index, and § 412.66(h), as adjusted by the capital construction cost index adopted by HCFA and applicable to the hospital.

G. Subpart H is amended as follows:

# Subpart H—Payments to Hospitals Under the Prospective Payment System

Section 412.113 is amended by revising paragraph (a) to read as follows:

# § 412.113 Payments determined on a reasonable cost basis.

(a) Capital costs. Payment for capital costs (as described in § 413.130 of this chapter) is determined on a reasonable cost basis for cost reporting periods beginning on or after October 1, 1983 and before October 1, 1987. During that period, the capital costs for each hospital must be determined consistently with the treatment of such costs for purposes of determining the hospital-specific portion of the hospital's prospective payment rate under §§ 412.70 through 412.73. For cost reporting periods beginning on or after October 1, 1987, capital costs are paid on a prospective basis as described in §§ 412.65 through 412.68. *

2. In § 412.125, the introductory language of the section is republished and paragraph (b) is revised to read as follows:

# § 412.125 Effect of change of ownership on payments under the prospective payment system.

When a hospital's ownership changes, as described in § 489.18 of this chapter, the following rules apply:

(b) Payment for capital costs (for reporting periods beginning before October 1, 1987) and bad debts, as described in §§ 412.113(a) and 412.115(a), respectively, is made to each owner or operator of the hospital (buyer and seller) in accordance with the principles of reasonable cost reimbursement.

H. In Subpart K, a new § 412.214 is added to read as follows:

Subpart K—Prospective Payment System for Hospitals Located in Puerto Rico

### § 412.214 Capital payments.

Subject to the blending percentages for the Federal rates described in § 412.204, capital payments for hospitals located in Puerto Rico are determined in the same manner, as described in §§ 412.65 through 412.68, as for other hospitals subject to the prospective payment system.

(Catalog of Federal Domestic Assistance Programs No. 13.773, Medicare-Hospital Insurance Program)

Dated: August 24, 1987.

### William L. Roper,

Administrator, Health Care Financing Administration.

Approved: August 27, 1987.

### Otis R. Bowen,

Secretary.

Note.—Appendices A-D will not appear in the Code of Federal Regulations.

### Appendix A—Federal Capital-Related Rates ¹

# TABLE 1.—FIFTY STATES AND DISTRICT OF COLUMBIA

	2279777007	Plant/fixed equipment		eable
11111	Urban	Rural	Urban	Rural
	180.03	160.86	122.39	92.72

¹ The rates reflect the reduction for capitalrelated costs of seven percent for cost reporting periods or discharges (as the case may be) occurring during FY 1988, as required under section 1886(g)(3)(A)(ii) of the Act, and an offset for budget neutrality as required under section 1886(g)(3)(C)(iii) of the Act for cost reporting periods occurring during FY 1988.

### TABLE 2.—PUERTO RICO 1

	Plant/fixed equipment		Moveable equipment	
.tstendy.a	Urban	Rural	Urban	Rural
Puerto Rico		167.55	38.29	3.88
National	nal 181.84		119.23	

¹ For hospitals located in Puerto Rico, the Federal rate is comprised of 75 percent of the Puerto Rico-specific adjusted capital prospective payment rate and 25 percent of the discharge-weighted average of the national urban and rural adjusted capital prospective payment rates.

# Appendix B—Construction Cost Indexes

# Table I.—Construction Cost Index for Urban Areas

for Urban Areas	
Urban area (constituent counties or county equivalents)	Con- struction cost index
Abilene, TX	.899
Taylor,TX Akron, OH	1.052
Portage OH	1.053
Summit, OH -	uniform.
Albany, GA	.814
Dougherty, GA	\$0 Wines
Lee, GA Albany-Schenectady-Troy, NY	1.190
Albany, NY	1.150
Greene, NY	dia a
Montgomery, NY	male.
Rensselaer, NY Saratoga, NY	January 2
Schenectady, NY	( nightop
Albuquerque, NM	.987
Bernalillo, NM	
Alexandria, LA	1.049
Rapides, LA	
Allentown-Bethlehem, PA NJ Warren, NJ	1.114
Carbon, PA	MAY PROT
Lehigh, PA	surelm
Northampton, PA	
Altoona, PA	1.464
Blair, PA Amarillo, TX	.917
Potter, TX	.017
Randall, TX	CHOCOHEU CONTROL
Anaheim-Santa Ana, CA	
Orange, CA Anchorage, AK	1.717
Anchorage, AK	4.75-0 (77)
Anderson, IN	.921
Madison, IN	augus never
Anderson, SC	.800
Ann Arbor, MI	1.166
Washtenaw, MI	
Anniston, AL	.830
Calhoun, AL Appleton-Oshkosh-Neenah, WI	.925
Calumet, WI	.925
Outagamie, WI	
Winnebago, WI	
Asheville, NC	.827
Buncombe, NC Athens, GA	.879
Clarke, GA	.073
Jackson, GA	
Madison, GA	
Oconee, GA Atlanta, GA	.844
Barrow, GA	.844
Butts, GA	
Cherokee, GA	
Clayton, GA	

# Table I.—Construction Cost Index for Urban Areas—Continued

for Orban Areas—Contin	lueu
Urban area (constituent counties or county equivalents)	Con- struction cost index
Cobb, GA	Standard
Coweta, GA	Ul Carlot
De Kalb, GA	and the same of
Douglas, GA	- Committee
Fayette, GA	UR WILLIAM
Forsyth, GA	
Fulton, GA Gwinnett, GA	ST TARRE
Henry, GA	Chantime
Newton, GA	and a second
Paulding, GA	
Rockdale, GA	
Spalding, GA	THE CO
Walton, GA	
Atlantic City, NJ	1.188
Atlantic, NJ	
Cape May, NJ	23.7
Augusta, GA-SC	.830
Columbia, GA McDuffie, GA	
Richmond, GA	1 1 2 2 2 2 2 2
Aiken, SC	FILMEN
Aurora-Elgin, IL	.917
Kane, IL	landar della
Kendall II	-
Austin, TX	.888
Hays, Tx	2
Travis, TX	
Williamson, TX	4 000
Bakersfield, CAKern, CA	1.082
Baltimore, MD	1.125
Anne Arundel, MD	4.447
Baltimore, MD	
Baltimore City, MD	
Carroll, MD	BELL
Harford, MD	12
Howard, MD	1601 F B
Queen Annes, MD	4 000
Bangor, MEPenobscot, ME	1.067
Baton Rouge, LA	.924
Ascension, LA	1041
East Baton Rouge, LA	
Livingston, LA	
West Baton Rouge, LA	
Battle Creek, MI	.985
Calhoun, MI Beaumont-Port Arthur, TX	
	.900
Hardin, TX	
Jefferson, TX Orange, TX	
Beaver County, PA	1 356
Beaver, PA	1.000
Beaver, PA Bellingham, WA	1.029
Whatcom WA	
Benton Harbor, MI	1.002
Berrien, MI	
Bergen-Passaic, NJ	1.353
Bergen, NJ Passaic, NJ	
Passaic, NJ	4.050
Billings, MT	1.056

# Table I.—Construction Cost Index for Urban Areas—Continued

### Con-Urban area (constituent counties struction or county equivalents) cost Yellowstone, MT Biloxi-Gulfport, MS. .958 Hancock, MS Harrison, MS Binghamton, NY. 1.268 Broome, NY Tioga, NY Birmingham, AL. .876 Blount, AL Jefferson, AL Saint Clair, AL Shelby, AL Walker, AL Bismarck, ND.. .957 Burleigh, ND Morton, ND Bloomington, IN 1.396 Monroe, IN Bloomington Normal, IL... .970 McLean, IL Boise City, ID .... .793 Ada, ID Boston-Lawrence-Salem-Lowell Brockton, MA .. 1.338 Essex, MA Middlesex, MA Norfolk, MA Plymouth, MA Suffolk, MA Boulder Longmont, CO ..... .882 Boulder, CO Bradenton, FL ... .760 Manatee, FL Brazoria, TX... .832 Brazoria, TX Bremerton, WA .. 1.069 Kitsap, WA Bridgeport-Stamford-Norwalk-Danbury, CT ..... 1.290 Fairfield, CT Brownsville Harlingen, TX..... .739 Cameron, TX Bryan-College Station, TX ... 1.012 Brazos, TX Buffalo, NY .... 1.352 Erie, NY Burlington, NC. .824 Alamance, NC Burlington, VT ..... .984 Chittenden, VT Grand Isle, VT Canton, OH .... .902 Carroll, OH Stark, OH Casper, WY .... .784 Natrona, WY Cedar Rapids, IA .858 Linn, IA Champaign-Urbana-Rantoul, IL... 1.243 Champaign, IL Charleston, SC ..... .921 Berkeley, SC Charleston, SC Dorchester, SC

# Table I.—Construction Cost Index for Urban Areas—Continued

for Urban Areas—Contin	nued
Urban area (constituent counties or county equivalents)	Con- struction cost index
Charleston, WV Kanawha, WV	1.155
Putnam, WV	
Charlotte-Gastonia-Rock Hill	400
NC-SC	.803
Cabarrus, NC Gaston, NC	link needs
Lincoln, NC	
Mecklenburg, NC	
Rowan, NC	
Union, NC	
York, SC Charlottesville, VA	1 242
Albermarle, VA	1.343
Charlottesville City, VA	
Fluvanna, VA	
Greene, VA Chattanooga, TN-GA	
Catoosa, GA	.868
Dade, GA	
Walker, GA	
Hamilton, TN	
Marion, TN Sequatchie, TN	
Cheyenne, WY	.976
Laramie, WY	1070
Chicago, IL	1.108
Cook, IL	Star St
Du Page, IL McHenry, IL	DEMERS
Chico, CA	.970
Butte, CA	
Cincinnati, OH-KY-IN	1.039
Dearborn, IN Boone, KY	Service Contract
Campbell, KY	
Kenton, KY	en 1037
Clermont, OH	1000
Hamilton, OH Warren, OH	COUNTY DIS
Clarksville-Hopkinsville, TN-KY	.904
Christian, KY	.904
Montgomery, TN	BILLET
Cleveland, OH	1.190
Cuyahoga, OH Geauga, OH	180
Lake, OH	1
Medina, OH	
Colorado Springs, CO	.847
El Paso, CO	
Columbia, MO	1.197
Columbia, SC	.844
Lexington, SC	- OLES
Richland, SC	THE STATE OF
Columbus, GA-AL  Russell, AL	.940
Chattanoochee, GA	5000
Muscogee, GA	ENVIOLE S
Columbus, OH	.978
Delaware, OH Fairfield, OH	THE REAL PROPERTY.
annotal Ott	THE PERSON NAMED IN

Franklin, OH

# Table I.—Construction Cost Index for Urban Areas—Continued

	Contin	lueu
	Urban area (constituent counties or county equivalents)	Con- struction cost
		index
The state of the s	Licking, OH Madison, OH Pickaway, OH Union, OH	
	Corpus Christi, TX	.904
	San Patricio, TX  Cumberland, MD-WV  Allegeny, MD	1.183
	Mineral, WV Dallas, TX Collin, TX	.861
	Dallas, TX Denton, TX	
	Ellis, TX Kaufman, TX Rockwall, TX	
	Danville, VA Danville, VA Pittsylvania, VA	.924
١	Davenport-Rock Island-Moline,	
	IA-ILScott, IA	1.016
	Henry, IL Rock Island, IL Dayton-Springfield, OH	
	Clark, OH Greene, OH	1.081
ı	Miami, OH	
	Montgomery, OH Daytona Beach, FLVolusia, FL	.903
	Decatur, IL	1.111
	Macon, IL Denver, CO Adams, CO	.932
	Arapahoe, CO Denver, CO	
	Douglas. CO	
	Jefferson, Co Des Moines. IA Dallas, IA	1.081
	Polk, IA Warren, IA	
	Detroit, MILapeer, MI	I.135
	Livingston, MI Macomb, MI Monroe, MI	
	Oakland, MI Saint Clair, MI	
	Wayne, MI Dothan, AL Dale, AL	.805
	Houston, AL Dubuque, IA	.982
	Dubuque, IA Duluth, MN-WI St. Louis, MN	1.063
	Douglas, WI Eau Claire, WI	.839

# Table I.—Construction Cost Index for Urban Areas—Continued

### Con-Urban area (constituent counties struction or county equivalents) cost index Chappewa, WI Eau Claire, WI El Paso, TX. .858 El Paso, TX Elkhart-Goshen, IN... 959 Elkhart, IN Elmira, NY .... 1.121 Chemung, NY Enid, OK .955 Garfield, OK Erie, PA... 1.223 Erie, PA Eugene-Springfield, OR 1.028 Lane, OR Evansville, IN-KY... 1.081 Posey, IN Vanderburgh, IN Warrick, IN Henderson, KY Fargo-Moorhead, ND-MN... .940 Clay, MN Cass, ND Fayetteville, NC. .785 Cumberland, NC Fayetteville-Springdale, AR .. .947 Washington, AR Flint, MI ..... .987 Genesee, MI Shiawassee, MI Florence, AL.... .861 Colbert, AL Lauderdale, AL Florence, SC .... .840 Florence, SC Fort Collins-Loveland, CO..... .926 Larimor, CO Fort Lauderdale-Hollywood-Pompano Beach, FL..... .878 Broward, FL Fort Myers-Cape Coral, FL ..... .761 Lee, FL Fort Pierce, FL ..... .900 Martin. FL St. Lucie, FL Fort Smith, AR-OK .770 Crawford, AR Sebastian, AR Sequoyah, OK Fort Walton Beach, FL ..... 1.008 Okaloosa, FL Fort Wayne, IN ... .964 Allen, IN De Kalb, IN Whitley, IN Fort Worth-Arlington, TX..... .883 Johnson, TX Parker, TX Tarrant, TX Fresno, CA 1.057 Fresno, CA Gadsden, AL... .813 Etowah, AL Gainesville, FL ... .989 Alachua, FL Bradford, FL

# Table I.—Construction Cost Index for Urban Areas—Continued

for Urban Areas—Continued			
Urban area (constituent counties or county equivalents)	Con- struction cost index		
Galveston-Texas City, TX	1.047		
Gary-Hammond, IN Lake, IN	1.050		
Porter, IN Glens Falls, NY Warren, NY	1.377		
Warren, NY Washington, NY Grand Forks, NDGrand Forks, ND	.943		
Grand Rapids, MI Kent, MI	.912		
Ottawa, MI Great Falls, MT Cascade, MT	1.069		
Greeley, CO	1		
Green Bay, WI Brown, WI	.859		
Greensboro-Winston-Salem-High Point, NC Davidson, NC	.850		
Davie, NC  Forsyth, NC	Pulse and		
Guilford, NG Randolph, NC	Service Co		
Stokes, NC Yadkin, NC			
Greenville-Spartanburg, SC Greenville, SC Pickens, SC	.804		
Spartanburg, SC Hagerstown, MD	.988		
Washington, MD Hamilton-Middletown, OH Butler, OH	.937		
Harrisburg-Lebanon-Carlisle, PA Cumberland, PA	1.109		
Dauphin, PA Lebanon, PA			
Perry, PA Hartford-Middletown-New Brit-	4 000		
ain-Bristol, CT Hartford, CT Litchfield, CT	1.220		
Middlesex, CT Tolland, CT			
Hickory, NC	826		
Burke, NC Catawba, NC	T-SOM		
Honolulu, HI Honolulu, HI	1.023		
Houma-Thibodaux, LALafourche, LA	1.201		
Terrebonne, LA Houston, TX	.858		
Fort Bend, TX Harris, TX Liberty, TX	Padage S		
Montgomery, TX			

Waller, TX

# Table I.—Construction Cost Index for Urban Areas—Continued

Urban area (constituent counties or county equivalents)  Huntington-Ashland, WV-KY-OH 1.029  Boyd, KY Carter, KY Creenup, KY Lawrence, OH Cabell, WV Wayne, WV Huntsville, AL		
or county equivalents)  Cost index  Huntington-Ashland, WV-KY-OH Boyd, KY Carter, KY Greenup, KY Lawrence, OH Cabell, WV Wayne, WV Huntsville, AL Madison, AL Indianapolis, IN Boone, IN Hamilton, IN Hancock, IN Hendricks, IN Johnson, IN Morgan, IN Shelby, IN Iowa City, IA Jackson, MI Jackson, MS Madison, MS Rankin, MS Madison, MS Rankin, MS Jackson, TN Jacksonville, FL Clay, FL Duval, FL Nassau, FL St. Johns, FL Jacksonville, NC Janesville-Beloit, WI Sonson, NS Carter, TN Hawkins, TN Sullivan, TN Unicoi, TN Washington, NA Boilt, IL Will, IL Joplin, MO Saleron, MO Kalmazoo, MI Jasper, MO Newaton, MO Kalmazoo, MI Jasper, MO Newaton, MI Jasper, MO Newaton, MO Kalmazoo, MI Kankake, II. L Manuelle, St. Johns Johnson Jington Jingto	- mile -	Con-
Huntington-Ashland, WV-KY-OH Boyd, KY Carter, KY Greenup, KY Lawrence, OH Cabell. WV Wayne, WV Huntsville, AL		
Huntington-Ashland, WV-KY-OH  Boyd, KY Carter, KY Greenup, KY Lawrence, OH Cabell. WV Wayne, WV Huntsville, AL	or county equivalents)	
Boyd, KY Carter, KY Greenup, KY Lawrence, OH Cabell, WV Wayne, WV Huntsville, AL		much
Boyd, KY Carter, KY Greenup, KY Lawrence, OH Cabell, WV Wayne, WV Huntsville, AL	Unstington Ashland WV VV OU	1.020
Carter, KY Greenup, KY Lawrence, OH Cabell, WV Wayne, WV Huntsville, AL		1.029
Greenup, KY Lawrence, OH Cabell, WV Wayne, WV Huntsville, AL		
Cabell, WV Wayne, WV Huntsville, AL		
Wayne, WV Huntsville, AL		
Huntsville, AL	Cabell, WV	
Madison, AL Indianapolis, IN	Wayne, WV	000
Indianapolis, IN		.006
Boone, IN Hamilton, IN Hancock, IN Hendricks, IN Johnson, IN Marion, IN Morgan, IN Shelby, IN Iowa City, IA	Indiananolis IN	988
Hancock, IN Hendricks, IN Johnson, IN Marion, IN Morgan, IN Shelby, IN Iowa City, IA Johnson, IA Jackson, MI Jackson, MI Jackson, MS Hinds, MS Madison, MS Rankin, MS Jackson, TN Jacksonville, FL Clay, FL Duval, FL Nassau, FL St. Johns, FL Jacksonville, NC Onslow, NC Janesville-Beloit, WI Rock, WI Jersey City, NJ Hudson, NJ Johnson City-Kingsport-Bristol, TN-VA Carter, TN Hawkins, TN Sullivan, TN Unicoi, TN Washington, TN Bristol City, VA Scott, VA Washington, VA Johnstown, PA Cambria, PA Somerset, PA Joliet, IL Will, IL Joplin, MO Jasper, MO Newton, MO Kalamazoo, MI Kankakee, IL Kankakee, IL Kankakee, IL Laton  1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370		
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Johnson, IN Marion, IN Morgan, IN Shelby, IN  Iowa City, IA Johnson, IA Jackson, MI Jackson, MI Jackson, MS Jackson, MS Madison, MS Rankin, MS Jackson, TN Jacksonville, FL Clay, FL Duval, FL Nassau, FL St. Johns, FL Jacksonville, NC Janesville-Beloit, WI Jersey City, NJ Hudson, NJ Johnson City-Kingsport-Bristol, TN-VA Carter, TN Hawkins, TN Sullivan, TN Unicoi, TN Washington, VA Johnstown, PA Cambria, PA Somerset, PA Joliet, IL Grundy, IL Will, IL Joplin, MO Jasper, MO Newton, MO Kalamazoo, MI Kankakee, IL Kankakee, IL Kankakee, IL Kankakee, II L L Johnstown, PA L L L L L L L L L L L L L L L L L L L		
Marion, IN Morgan, IN Shelby, IN Iowa City, IA. 1.370 Johnson, IA Jackson, MI 1.096 Jackson, MI 930 Hinds, MS Madison, MS Rankin, MS Jackson, TN. 937 Madison, TN Jacksonville, FL 874 Clay, FL Duval, FL Nassau, FL St. Johns, FL Jacksonville, NC. 1.115 Onslow, NC Janesville-Beloit, WI 932 Rock, WI Jersey City, NJ 1.400 Hudson, NJ Johnson City-Kingsport-Bristol, TN-VA 1.004 Carter, TN Hawkins, TN Sullivan, TN Unicoi, TN Washington, TN Bristol City, VA Scott, VA Washington, VA Johnstown, PA 1.199 Cambria, PA Somerset, PA Joliet, IL 877 Grundy, IL Will, IL Joplin, MO 941 Jasper, MO Newton, MO Kalamazoo, MI Kalamazoo, MI Kankakee, IL Kankakee, IL Kankakee, IL Kankakee, II. 1.119		
Morgan, IN Shelby, IN Iowa City, IA Johnson, IA Jackson, MI Jackson, MI Jackson, MS Madison, MS Rankin, MS Jackson, TN Madison, TN Jacksonville, FL Clay, FL Duval, FL St. Johns, FL Jacksonville, NC Janesville-Beloit, WI Rock, WI Jersey City, NJ Hudson, NJ Johnson City-Kingsport-Bristol, TN-VA Carter, TN Hawkins, TN Sullivan, TN Unicoi, TN Washington, VA Johnstown, PA Cambria, PA Somerset, PA Joliet, IL Will, IL Joplin, MO Jasper, MO Newton, MO Kalamazoo, MI Kankakee, IL Ling 1.096 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370 1.370	A CONSISSION CONTRACT	
Shelby, IN Iowa City, IA Johnson, IA Jackson, MI Jackson, MI Jackson, MS Hinds, MS Madison, MS Rankin, MS Jackson, TN Jacksonville, FL Clay, FL Duval, FL Nassau, FL St. Johns, FL Jacksonville, NC Janesville-Beloit, WI Rock, WI Jersey City, NJ Hudson, NJ Johnson City-Kingsport-Bristol, TN-VA Carter, TN Hawkins, TN Sullivan, TN Unicoi, TN Washington, VA Johnstown, PA Cambria, PA Somerset, PA Joliet, IL Grundy, IL Will, IL Joplin, MO Jasper, MO Newton, MO Kalamazoo, MI Kankakee, IL Kankakee, IL Kankakee, IL Kankakee, II L 1.098 1.370 1.098 1.098 1.098 1.098 1.098 1.098 1.098 1.098 1.098 1.098 1.098 1.098 1.098 1.098 1.098 1.098 1.098 1.119 1.119 1.119		BUT FUE
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Johnson, IA Jackson, MI Jackson, MS Jackson, MS Hinds, MS Madison, MS Rankin, MS Jackson, TN Jacksonville, FL Clay, FL Duval, FL Nassau, FL St. Johns, FL Jacksonville, NC Janesville-Beloit, WI Rock, WI Jersey City, NJ Hudson, NJ Johnson City-Kingsport-Bristol, TN-VA Carter, TN Hawkins, TN Sullivan, TN Unicoi, TN Washington, TN Bristol City, VA Scott, VA Washington, VA Johnstown, PA Cambria, PA Somerset, PA Joliet, IL Will, IL Joplin, MO Jasper, MO Newton, MO Kalamazoo, MI Kalamazoo, MI Kalamazoo, MI Kalamazoo, MI Kankakee, IL Kankakee, IL Kankakee, II. L1090  1.093  1.096 1.096 1.096 1.096 1.096 1.097 1.098 1.098 1.098 1.098 1.098 1.098 1.098 1.098 1.098 1.098 1.119 1.098 1.098 1.098 1.098 1.119 1.098 1.098 1.098 1.111	Iowa City, IA	1.370
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Jackson, MS Hinds, MS Madison, MS Rankin, MS Jackson, TN Jacksonville, FL Clay, FL Duval, FL Nassau, FL St. Johns, FL Jacksonville, NC Onslow, NC Janesville-Beloit, WI Jersey City, NJ Hudson, NJ Johnson City-Kingsport-Bristol, TN-VA Carter, TN Hawkins, TN Sullivan, TN Unicoi, TN Washington, TN Bristol City, VA Scott, VA Washington, VA Johnstown, PA Cambria, PA Somerset, PA Joliet, IL Will, IL Joplin, MO Newton, MO Kalamazoo, MI Kalamazoo, MI Kankakee, IL Kankakee IL Kankakee II.	Jackson, MI	1.096
Hinds, MS Madison, MS Rankin, MS Jackson, TN Madison, TN Jacksonville, FL Clay, FL Duval, FL Nassau, FL St. Johns, FL Jacksonville, NC Janesville-Beloit, WI Rock, WI Jersey City, NJ Hudson, NJ Johnson City-Kingsport-Bristol, TN-VA Carter, TN Hawkins, TN Sullivan, TN Unicoi, TN Washington, TN Bristol City, VA Scott, VA Washington, VA Johnstown, PA Cambria, PA Somerset, PA Joliet, IL Will, IL Joplin, MO Jasper, MO Newton, MO Kalamazoo, MI Kalamazoo, MI Kankakee, IL Kankakee IL Kankakee II.  1.937  4874  874  875  876  877  877  877  877  877  877	Jackson, MI	000
Madison, MS Rankin, MS Jackson, TN Madison, TN Jacksonville, FL Clay, FL Duval, FL Nassau, FL St. Johns, FL Jacksonville, NC Janesville-Beloit, WI Jersey City, NJ Hudson, NJ Johnson City-Kingsport-Bristol, TN-VA Carter, TN Hawkins, TN Sullivan, TN Unicoi, TN Washington, TN Bristol City, VA Scott, VA Washington, VA Johnstown, PA Cambria, PA Somerset, PA Joliet, IL Will, IL Joplin, MO Jasper, MO Newton, MO Kalamazoo, MI Kankakee, IL Kankakee, IL Kankakee, II. 1.874  874  875  1.119  876  877  877  941  1.119		.930
Rankin, MS Jackson, TN Madison, TN Jacksonville, FL Clay, FL Duval, FL Nassau, FL St. Johns, FL Jacksonville, NC Onslow, NC Janesville-Beloit, WI Rock, WI Jersey City, NJ Hudson, NJ Johnson City-Kingsport-Bristol, TN-VA Carter, TN Hawkins, TN Sullivan, TN Unicoi, TN Washington, TN Bristol City, VA Scott, VA Washington, VA Johnstown, PA Cambria, PA Somerset, PA Joliet, IL Will, IL Joplin, MO Jasper, MO Newton, MO Kalamazoo, MI Kalamazoo, MI Kankakee, IL Kankakee, IL Kankakee, II. 1.874  874  874  1.115  875  876  877  877  877  877  877  87	Madison, MS	FILE .
Jackson, TN.  Madison, TN  Jacksonville, FL.  Clay, FL  Duval, FL  Nassau, FL  St. Johns, FL  Jacksonville, NC.  Onslow, NC  Janesville-Beloit, WI  Rock, WI  Jersey City, NJ  Hudson, NJ  Johnson City-Kingsport-Bristol,  TN-VA  Carter, TN  Hawkins, TN  Sullivan, TN  Unicoi, TN  Washington, TN  Bristol City, VA  Scott, VA  Washington, VA  Johnstown, PA  Cambria, PA  Somerset, PA  Joliet, IL  Will, IL  Joplin, MO  Jasper, MO  Newton, MO  Kalamazoo, MI  Kankakee, IL  Kankakee, IL  Kankakee, II.  1.874  .874  .874  .874  .874  .875  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .877  .878  .877  .877  .877  .878  .877  .877  .878  .878  .877  .877  .878  .877  .878  .878  .877  .878  .878  .877  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878  .878	Rankin, MS	
Jacksonville, FL	Jackson, TN	.937
Clay, FL Duval, FL Nassau, FL St. Johns, FL Jacksonville, NC	Madison, TN	
Duval, FL Nassau, FL St. Johns, FL Jacksonville, NC		.874
Nassau, FL St. Johns, FL Jacksonville, NC		III.
St. Johns, FL Jacksonville, NC		CONT.
Onslow, NC Janesville-Beloit, WI		Comment.
Janesville-Beloit, WI		1.115
Rock, WI Jersey City, NJ	Onslow, NC	000
Jersey City, NJ Hudson, NJ Johnson City-Kingsport-Bristol, TN-VA Carter, TN Hawkins, TN Sullivan, TN Unicoi, TN Washington, TN Bristol City, VA Scott, VA Washington, VA Johnstown, PA Cambria, PA Somerset, PA Joliet, IL Grundy, IL Will, IL Joplin, MO Newton, MO Kalamazoo, MI Kankakee, IL Kankakee IL  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.400  1.		.932
Hudson, NJ Johnson City-Kingsport-Bristol, TN-VA		1.400
TN-VA Carter, TN Hawkins, TN Sullivan, TN Unicoi, TN Washington, TN Bristol City, VA Scott, VA Washington, VA Johnstown, PA Cambria, PA Somerset, PA Joliet, IL Will, IL Joplin, MO Jasper, MO Newton, MO Kalamazoo, MI Kankakee, IL Kankakee, IL Kankakee, IL Kankakee, II.  1.004 1.004 1.004 1.004 1.119 1.119	Hudson, NJ	THE REAL PROPERTY.
Carter, TN Hawkins, TN Sullivan, TN Unicoi, TN Washington, TN Bristol City, VA Scott, VA Washington, VA Johnstown, PA Cambria, PA Somerset, PA Joliet, IL Will, IL Joplin, MO Jasper, MO Newton, MO Kalamazoo, MI Kankakee, IL Kankakee II.  Sullivan Hawkins, TN Hawkington, TN Haw	Johnson City-Kingsport-Bristol,	
Hawkins, TN Sullivan, TN Unicoi, TN Unicoi, TN Washington, TN Bristol City, VA Scott, VA Washington, VA Johnstown, PA Cambria, PA Somerset, PA Joliet, IL Will, IL Joplin, MO Jasper, MO Newton, MO Kalamazoo, MI Kankakee, IL Kankakee, IL Kankakee, IL Kankakee, II.		1.004
Sullivan, TN Unicoi, TN Washington, TN Bristol City, VA Scott, VA Washington, VA Johnstown, PA Cambria, PA Somerset, PA Joliet, IL Will, IL Joplin, MO Jasper, MO Newton, MO Kalamazoo, MI Kankakee, IL Kankakee IL Kankakee II.		
Unicoi, TN Washington, TN Bristol City, VA Scott, VA Washington, VA Johnstown, PA Cambria, PA Somerset, PA Joliet, II		D MANNET
Washington, TN Bristol City, VA Scott, VA Washington, VA Johnstown, PA Cambria, PA Somerset, PA Joliet, IL Grundy, IL Will, IL Joplin, MO Newton, MO Kalamazoo, MI Kankakee, IL Kankakee IL Kankakee II.		
Scott, VA Washington, VA Johnstown, PA Cambria, PA Somerset, PA Joliet, IL Will, IL Joplin, MO Jasper, MO Newton, MO Kalamazoo, MI Kankakee, IL Kankakee IL Kankakee IL Kankakee II.		The state of the s
Washington, VA  Johnstown, PA		- Back
Cambria, PA Somerset, PA Joliet, IL	Scott, VA	MITTER
Cambria, PA Somerset, PA  Joliet, IL	Washington, VA	1 100
Somerset, PA   Some	Cambria. PA	4,100
Joliet, IL	Somerset, PA	1
Grundy, IL Will, IL Joplin, MO Jasper, MO Newton, MO Kalamazoo, MI Kalamazoo, MI Kankakee, IL Kankakee IL Kankakee IL	Joliet, IL	.877
Joplin, MO	Grundy, IL	
Jasper, MO Newton, MO Kalamazoo, MI Kalamazoo, MI Kankakee, IL Kankakee II.		0.11
Newton, MO		
Kalamazoo, MI  Kalamazoo, MI  Kankakee, IL  Kankakee II.  1.119	Newton MO	place 7
Kalamazoo, MI Kankakee, IL	Kalamazoo, MI	1.117
Kankakee II. 1.119	Kalamazoo, MI	The state of the s
Kankakee, IL Kansas City, KS-MO	Kankakee, IL	1.119
Kansas Chy, Ko-WO	Kankakee, IL	913
	Kansas City, Ko WO	1 1000

Table	I.—Construction Cost Index
for	Urban Areas—Continued

for Urban Areas—Contin	for Urban Areas—Contin	
Urban area (constituent counties or county equivalents)	Con- struction cost index	Urban area (constituent counties or county equivalents)
Johnson, KS		Allen, OH
Leavenworth, KS	AMLANK!	Auglaize, OH
Miami, KS	Daniti Ell	Lincoln, NE
Wyandotte, KS		Lancaster, NE
Cass, MO	State of the last	Little Rock-North Little Rock, AR
Clay, MO		Faulkner, AR
Jackson, MO		Lonoke, AR
Lafayette, MO Platte, MO	daniel - 1	Pulaski, AR
Ray, MO	HET S	Saline, AR Longview-Marshall, TX
Kenosha, WI	.872	Gregg, TX
Kenosha, WI	.072	Harrison, TX
Killeen-Temple, TX	.910	Lorain-Elyria, OH
Bell, TX		Lorain, OH
Coryell, TX		Los Angeles-Long Beach, CA
Knoxville, TN	.873	Los Angeles, CA
Anderson, TN		Louisville, KY-IN
Blount, TN	TO SERVICE TO	Clark, IN
Grainger, TN Jefferson, TN	OF OWNER, DE	Floyd, IN
Knox, TN	THE LEGICAL DE	Harrison, IN Bullitt, KY
Sevier, TN		Jefferson, KY
Union, TN	NEWS CO.	Oldham, KY
Kokomo, IN	.885	Shelby, KY
Howard, IN	SEE SEE	Lubbock, TX
Tipton, IN	and the	Lubbock, TX
LaCrosse, WI	.970	Lynchburg, VA
LaCrosse, WI	0.00	Amherst, VA
Lafayette, LA Lafayette, LA	.958	Campbell, VA
St. Martin, LA		Lynchburg City, VA
Lafayette, IN	1.116	Macon-Warner Robins, GA Bibb, GA
Tippecanoe, IN	4.110	Houston, GA
Lake Charles, LA	.929	Jones, GA
Calcasieu, LA	THE PARTY	Peach, GA
Lake County, IL	1.035	Madison, WI
Lake, IL	MODES	Dane, WI
Lakeland-Winter Haven, FL Polk, FL	.788	Manchester-Nashua, NH
Lancaster, PA	.977	Hillsborough, NH Merrimack, NH
Lancaster, PA	.3//	Mansfield, OH
Lansing-East Lansing, MI	1.108	Richland, OH
Clinton, MI	EDITE S	McAllen-Edinburg-Mission, TX
Eaton, MI	III.	Hidalgo, TX
Ingham, MI	P (Shirt	Medford, OR
Laredo, TX	.728	Jackson, OR
Las Cruces, NM	.845	Melbourne-Titusville, FL
Dona Ana, NM	640.	Brevard, FL Memphis, TN-AR-MS
Las Vegas, NV	.949	Crittenden, AR
Clark, NV	S D	De Soto, MS
Lawrence, KS	1.118	Shelby, TN
Douglas, KS		Tipton, TN
Lawton, OK	.997	Merced, CA
Comanche, OK Lewiston-Auburn, ME	orn	Merced, CA
Androscoggin, ME	.952	Miami-Hialeah, FL
Lexington-Fayette, KY	1.063	Dade, FL Middlesex-Somerset-Hunterdon,
Bourbon, KY	2,000	NJ
Clark, KY	THE REST	Hunterdon, NJ
Fayette, KY	HEIDER!	Middlesex, NJ
Jessamine, KY		Somerset, NJ
Scott, KY Woodford, KY	Will be seen to be see	Midland, TX
Woodford, KY Lima, OH	.943	Midland, TX
011	.843	Milwaukee, WI

# Table I.—Construction Cost Index

	Con-
Urban area (constituent counties	struction
or county equivalents)	cost
	index
Alles OU	Solie .
Allen, OH Auglaize, OH	The same
Lincoln, NE	1.066
Lancaster, NE	
Little Rock-North Little Rock, AR	.925
Faulkner, AR	
Lonoke, AR	
Pulaski, AR Saline, AR	100000
Longview-Marshall, TX	.746
Gregg, TX	., 10
Harrison, TX	
Lorain-Elyria, OH	1.048
Lorain, OH	Name of the last
Los Angeles-Long Beach, CA	1.065
Los Angeles, CA Louisville, KY-IN	.983
Clark, IN	.903
Floyd, IN	HEESE.
Harrison, IN	winn.
Bullitt, KY	neven n
Jefferson, KY	THE PARTY
Oldham, KY	Co office
Shelby, KY Lubbock, TX	.911
Lubbock, TX	.911
Lynchburg, VA	.900
Amherst, VA	18-110
Campbell, VA	1343/2
Lynchburg City, VA Macon-Warner Robins, GA	000
Bibb, GA	.809
Houston, GA	
Jones, GA	
Peach, GA	
Madison, WI	1.044
Dane, WI Manchester-Nashua, NH	4.004
Hillsborough, NH	1.004
Merrimack, NH	
Mansfield, OH	.875
Richland, OH	
McAllen-Edinburg-Mission, TX	.762
Hidalgo, TX Medford, OR	.975
Jackson, OR	.9/5
Melbourne-Titusville, FL	.839
Brevard, FL	-
Memphis, TN-AR-MS	.877
Crittenden, AR	
De Soto, MS Shelby, TN	
Tipton, TN	
Merced, CA	1.092
Merced, CA	1002
Miami-Hialeah, FL	.961
Dade, FL	
Middlesex-Somerset-Hunterdon,	1 2322
NJ Hunterdon, NJ	1.191
Middlesex, NJ	
Somerset, NJ	
Midland, TX	.862
Midland, TX	

# Table I.—Construction Cost Index for Urban Areas-Continued

Urban area (constituent counties or county equivalents)	Con- struction cost index
Milwaukee MI	Allen .
Milwaukee, WI Ozaukee, WI	THE RESERVE
Washington, WI	THE REAL PROPERTY.
Waukesha, WI	
Minneapolis-St. Paul, MN-WI	040
Anoka, MN	.943
Carver, MN	A THE REAL PROPERTY.
Chisago, MN Dakota, MN	
Hennepin, MN	APPROPRIEST OF
Isanti, MN	
Ramsey, MN	
Scott, MN	
Washington, MN	
Wright, MN	
St. Croix, WI	
Mobile, AL	.906
Baldwin, AL	,500
Mobile AL	
Mobile, AL Modesto, CA	1.058
Stanislaus, CA	1.000
Monmouth-Ocean, NJ	1 093
Monmouth, NI	1,000
Ocean, NJ	
Monroe, LA	.934
Ouachita, LA	
Montgomery, AL	.824
Autauga, AL	Service .
Elmore, AL	
Montgomery, AL	
Muncie, IN	1.006
Delaware, IN	
Muskegon, MI	.970
Muskegon, MI	
Naples, FL	.879
Collier, FL	
Nashville, TN	.875
Cheatham, TN	
Davidson, TN	
Dickson, TN	
Robertson, TN	
Rutherford, TN	
Sumner, TN	
Williamson, TN	
Wilson, TN	
Nassau-Suffolk, NY	1.382
Nassau, NY	
Suffolk, NY	
New Bedford-Fall River-Attleboro,	
MA	1.248
Bristol, MA	
New Haven-Waterbury-Meriden,	200
CT	1.199
New Haven, CT	2.244
New London-Norwich, CT	1.185
New London, CT	
New Orleans, LA	1.052
Tofferen I A	
Jefferson, LA	
Orleans, LA	
Orleans, LA St. Bernard, LA	
Orleans, LA St. Bernard, LA St. Charles, LA	
Orleans, LA St. Bernard, LA	

# Table I.—Construction Cost Index for Urban Areas—Continued

### Con-Urban area (constituent counties struction or county equivalents) cost index Bronx, NY Kings, NY New York City, NY Putnam, NY Queens, NY Richmond, NY Rockland, NY Westchester, NY Newark, NJ. 1.421 Essex, NJ Morris, NJ Sussex, NJ Union, NJ Niagara Falls, NY 1.190 Niagara, NY Norfolk-Virginia Beach-Newport News, VA ..... .957 Chesapeake City, VA Gloucester, VA Hampton City, VA James City Co., VA Newport News City, VA Norfolk City, VA Poquoson, VA Portsmouth City, VA Suffolk City, VA Virginia Beach City, VA Williamsburg City, VA York, VA Oakland, CA... 1.069 Alameda, CA Contra Costa, CA Ocala, FL ..... .755 Marion, FL Odessa, TX.. .969 Ector, TX Oklahoma City, OK. .865 Canadian, OK Cleveland, OK Logan, OK McClain, OK Oklahoma, OK Pottawatomie, OK Olympia, WA. 1.045 Thurston, WA Omaha, NE-IA. .958 Pottawattamie, IA Douglas, NE Sarpy, NE Washington, NE Orange County, NY. 1.382 Orange, NY Orlando, FL .... .858 Orange, FL Osceola, FL Seminole, FL Owensboro, KY... .959 Daviess, KY Oxnard-Ventura, CA 1.148 Ventura, CA Panama City, FL. .876 Bay. FL Parkersburg-Marietta, WV-OH... 1.075 Washington, OH Wood, WV Pascagoula, MS

# Table I.—Construction Cost Index for Urban Areas—Continued

	for Urban Areas—Contin	ued
on- ction ost dex	Urban area (constituent counties or county equivalents)	Con- struction cost index
	Jackson, MS Pensacola, FL Escambia, FL	.842
	Santa Rosa, FL Peoria, IL Peoria, IL	.968
1.421	Tazewell, IL Woodford, IL Philadelphia, PA-NJ Burlington, NJ	1.169
	Camden, NJ Gloucester, NJ Bucks, PA	
1.190	Chester, PA Delaware, PA Montgomery, PA	
.957	Philadelphia, PA Phoenix, AZ Maricopa, AZ	.959
	Pine Bluff, AR Jefferson, AR	.943
	Pittsburgh, PA	1.311
	Washington, PA Westmoreland, PA Pittsfield, MA	1.502
1.069	Berkshire, MA Portland, ME Cumberland, ME	.992
THE REAL PROPERTY.	Sagadahoc, ME York, ME	
.755	Portland, ORClackamas, OR Multnomah, OR	.989
.865	Washington, OR Yamhill, OR Portsmouth-Dover-Rochester, NH	1.050
	Rockingham, NH Strafford, NH Poughkeepsie, NY Dutchess, NY	1.193
1.045	Providence-Pawtucket- Woonsocket, RI Bristol, RI	1.101
.958	Kent, RI Newport, RI Providence, RI Washington, RI	
1.382	Provo-Orem, UT	Eastern Town
.858	Pueblo, CO Puerto Rico	1.000
.959	Racine, WIRacine, WI Raleigh-Durham, NC	1.088
1.148	Durham, NC Franklin, NC	200
.876	Orange, NC Wake, NC Rapid City, SD	1.112
L.075	Pennington, SD Reading, PA	
.757	Berks, PA Redding, CA	.944

# Table I.—Construction Cost Index for Urban Areas—Continued

	Con-
Urban area (constituent counties	
or county equivalents)	cost
Capacity September 1999	index
Shasta, CA	
Reno, NV	1.021
Washoe, NV	THE PARTY
Richland-Kennewick, WA	1.002
Benton, WA	Topper .
Franklin, WA Richmond-Petersburg, VA	1.011
Charles City Co., VA	1.011
Chesterfield, VA	Aleman .
Colonial Heights City, VA	NOT THE REAL PROPERTY.
Dinwiddie, VA Goochland, VA	
Goochland, VA	consit .
Hanover, VA Henrico, VA	THE PARTY IN
Hopewell City, VA	
New Kent, VA	ADDRESS.
Petersburg City, VA	
Powhatan, VA	i=mill:
Prince George, VA	Marie S
Richmond City, VA Riverside-San Bernardino, CA	1.073
Riverside, CA	1.0/3
San Bernardino, CA	
Roanoke, VA	.983
Botelourt, VA	Total Park
Roanoke, VA	
Roanoke, VA Roanoke City, VA Salem City, VA	Cit
Salem City, VA Rochester, MN	1.358
Olmsted MN	1.000
Rochester, NY	1.250
Livingston, NY	MAN WE
Monroe, NY	
Ontario, NY	The state of
Orleans, NY Wayne, NY	100
Rockford, IL	.899
Doors II	
Winnebago, IL	
Sacramento, CA	1.202
Eldorado, CA	
Placer, CA Sacramento, CA	
Yolo, CA	
Saginaw-Bay City-Midland, MI	1.013
Bay, MI	
Midland, MI	
Saginaw, MI St. Cloud, MN	nes
Ronton MN	.804
Sherburne, MN	
Benton, MN Sherburne, MN Stearns, MN	
St. Joseph, MO	1.035
Buchanan, MO	
St. Louis, MO-IL	1.143
Clinton, IL Jersey, IL	
Madison, IL	
Monroe, IL	
St. Clair, IL	
Franklin, MO	
Jefferson, MO	
St. Charles, MO	
St. Louis, MO St. Louis City, MO	
Salem, OR	000

# Table I.—Construction Cost Index for Urban Areas—Continued

### Con-Urban area (constituent counties struction or county equivalents) cost index Marion, OR Polk, OR Salinas-Seaside-Monterey, CA ...... 1.091 Monterey, CA Salt Lake City-Ogden, UT ..... .871 Davis, UT Salt Lake, UT Weber, UT San Angelo, TX. .909 Tom Green, TX San Antonio, TX .... .921 Bexar, TX Comal, TX Guadalupe, TX San Diego, CA... 1.043 San Diego, CA San Francisco, CA... 1.043 Marin, CA San Francisco, CA San Mateo, CA San Jose, CA .... 1.028 Santa Clara, CA Santa Barbara-Santa Maria Lompoc, CA ...... 1.036 Santa Barbara, CA Santa Cruz, CA ..... .950 Santa Cruz, CA Santa Fe, NM .... 1.086 Los Alamos, NM Santa Fe, NM Santa Rosa-Petaluma, CA ..... 1.134 Sonoma, CA Sarasota, FL.. 804 Sarasota, FL Savannah, GA ... 1.028 Chatham, GA Effingham, GA Scranton-Wilkes-Barre, PA ..... 1.110 Columbia, PA Lackawanna, PA Luzerne, PA Monroe, PA Wyoming, PA Seattle, WA.... 1.085 King, WA Snohomish, WA Sharon, PA .... 1.144 Mercer, PA Sheboygan, WI., 1.040 Sheboygan, WI Sherman-Denison, TX.... .949 Grayson, TX Shreveport, LA .... .936 Bossier, LA Caddo, LA Sioux City, IA-NE .. 1.020 Woodbury, IA Dakota, NE Sioux Falls, SD .. 1.202 Minnehaha, SD South Bend-Mishawaka, IN..... 1.143 St. Joseph, IN Spokane, WA ... .971 Spokane, WA

# Table I.—Construction Cost Index for Urban Areas—Continued

	1
Urban area (constituent counties or county equivalents)	Con- struction cost index
Springfield, IL	1.011
Sangamon, IL	AUTO OF
Springfield, MA	1.233
Hampden, MA	
Hampshire, MA Springfield, MO	.912
Christian, MO	.912
Greene, MO	lennite !
State College, PA	1.035
Centre, PA Steubenville-Weirton, OH-WV	1.167
Jefferson, OH	1.10/
Brooke, WV	130.0
Hancock, WV	100
Stockton, CA	1.060
Syracuse, NY	1.312
Madison, NY	COP.
Onondaga, NY	m wood
Oswego, NY Tacoma, WA	1.038
Pierce, WA	1.000
Tallahassee, FL	.989
Gadsden, FL	DOMESTIC BY
Leon, FL Tampa-St. Petersburg-Clearwater,	Shille
FL	.890
Hernando, FL	MA A
Hillsborough, FL	N US
Pasco, FL Pinellas, FL	rimbenden
Terre Haute, IN	1.040
Clay, IN	
Vigo, IN	DICTAL PL
Texarkana-TX-Texarkana, AR Miller, AR	.754
Bowie, TX	MA I
Toledo, OH	1.044
Fulton, OH	- Statemen
Lucas, OH Wood, OH	AND STATE
Topeka, KS	.996
Shawnee, KS	CON DI
Trenton, NJ	1.226
Mercer, NJ Tucson, AZ	.931
Pima, AZ	
Tulsa, OK	.911
Creeks, OK	TO HEALTH STATE
Osage, OK Rogers, OK	
Wagoner, OK	Figure
Tuscaloosa, AL	1.003
Tuscaloosa, AL	000
Tyler, TXSmith, TX	.922
Utica-Rome, NY	1.256
Herkimer, NY	I C. AL
Oneida, NY	74.7
Vallejo-Fairfield-Napa, CA Napa, CA	1.118
Solano, CA	
Vancouver, WA	.819

Clark, WA

# Table I.—Construction Cost Index for Urban Areas—Continued

101 Otban Areas—Contin	ueu
Urban area (constituent counties or county equivalents)	cost
	index
Victoria, TX	.842
Vineland-Millville-Bridgeton, NJ Cumberland, NJ	1.099
Visalia-Tulare-Porterville, CA	1.017
Waco, TX	.840
Washington, DC-MD-VA District of Columbia, DC	1.155
Calvert, MD Charles, MD	
Frederick, MD	
Montgomery, MD	BURNESS TO SERVICE
Prince Georges, MD	BY THE
Alexandria City, VA Arlington, VA	
Fairfax, VA	
Fairfax City, VA	Selement .
Falls Church City, VA	a built have been
Loudoun, VA Manassas City, VA	ALTERNATION OF THE PARTY OF THE
Manassas Park City, VA	Beldie
Prince William, VA	E PAR
Stafford, VA	A STATE OF THE PARTY OF THE PAR
Waterloo-Cedar Falls, IA	.911
Bremer, IA	
Wausau, WI	.889
Marathon, WI	
West Palm Beach-Boca Raton- Delray Beach, FL	.944
Palm Beach, FL Wheeling, WV-OH	4.050
Belmont, OH	1.052
Marshall, WV	
Ohio, WV	
Wichita, KS	.872
Butler, KS	
Harvey, KS Sedgwick, KS	
Wichita Falls, TX	.929
Wichita, TX	
Williamsport, PA	1.063
Lycoming, PA Wilmington, DE-NJ-MD	1 231
New Castle, DE	1.601
Cacil MD	
Salem, NJ	
Wilmington, NC	.851
Worcester-Fitchburg-Leominster,	
MAWorcester, MA	1.252
Yakima, WA	.975
Yakima, WA	
York, PA	.984
Adams, PA	
York, PA	
Youngstown-Warren, OH Mahoning, OH	.981
Trumbull, OH Yuba City, CA	10 TAX 10
Tuba City, CA	.910

# Table I.—Construction Cost Index for Urban Areas-Continued

Urban area (constituent counties or county equivalents)	Con- struction cost index
Sutter, CA Yuba, CA	A STATE OF

# TABLE II.—CONSTRUCTION COST INDEX FOR RURAL AREAS

	Nonurban area	Con- struction cost index
Alahama		.787
		1.582
		1.000
		.749
		1.034
	L	100000000000000000000000000000000000000
Connecti	cut	.939
	)	1.040
120000000000000000000000000000000000000		.817
		.930
		.978
		.919
		.880
		.814
		.921
	······	941
		1.037
Maryland		1.180
	usetts	1.169
		.962
	a	.918
	pi	.834
		.847
		.917
	1	.772
Nevada		.966
New Han	npshire	1.069
New Jers	sey 1	
New Mex	dico	.893
New York	<	1.224
North Ca	rolina	.788
North Da	kota	.942
Ohio		.884
Oklahom	a	.828
		.931
Pennsylv	ania	1.177
Puerto R	ico	1.000
Rhode Is	land 1	
South Ca	rolina	.774
	kota	.898
Tennesse	90	.766
		.783
Utah		.869
		1.038
		.912
	on	1.072
	jinia	1.031
	ń	.864
		.958
_100		and the same

¹ All counties within the State are classified urban.

# Appendix C-List of Plant and Fixed Equipment, and Moveable Equipment 1

A. Land Movements

Bumpers Culverts

Fencing

a. Brick or stone

b. Chain link

c. Wire

d. Wood

Flagpole

Heated pavement

Lawn sprinkler system

Parking lot gate

Parking lot, open walls

Paving (including roadways, walks, and parking)

a. Asphalt

b. Concrete

c. Gravel

Retaining wall

Shrubs, lawns, trees

Snow melting system

Turf, artificial

Underground sewer and water lines Yard lighting

B. Buildings

Boiler house

Garage

a. Masonry

b. Wood frame

Masonry, reinforced concrete frame Masonry, steel frame, fireproofed

Masonry, steel frame, not fireproofed

Masonry, wood frame

Reinforced concrete, common design Residence

a. Masonry

b. Wood frame

Storage building

a. Masonry

b. Wood frame

Building, componentized parts

a. Automatic door

b. Canopies

c. Ceiling finishes

d. Computer flooring

e. Cubicle track

f. Designation signs

g. Drapery track

h. Floor finishes

i. Folding partitions

j. Interior finish

k. Loading docks

1. Overhead door

m. Partitions, interior

n. Roof covering

o. Storefront construction

p. Toilet partitions

q. Wall paint

r. Wallpaper

Adapted from the list published by the American Hospital Association, 1987 edition. Multilevel parking structure, masonry

II. Fixed Equipment

A. Building Services Equipment

Boiler smokestack, metal

Clean air equipment

Clock system, central

Doctors' in-and-out register

Electric lighting and power

a. Feed wiring

b. Conduit and wiring

c. Fixtures

d. Transformer

e. Switch gear

Elevator

a. Dumbwaiter

b. Freight

c. Passenger, high-speed automatic

d. Passenger, other

Emergency light system

Escalator

Fire alarm system, door closing devices

Heating, ventilating, and air

conditioning system

Air conditioning system, all equipment and units

a. Large-over 20 tons

b. Medium-5-15 tons

c. Small-under 5 tons

d. Boiler

e. Compressor, air

f. Condensate tank

g. Condenser

h. Controls

i. Cooler and dehumidifier

j. Cooling tower

(1) Metal

(2) Wood k. Duct work

I. Fan, air handling and ventilating

m. Furnace, domestic type

n. Incinerator, indoor

o. Oil storage tank

p. Piping

q. Precipitator

r. Pump

s. Radiator, cast iron

t. Radiator, finned tube

u. Solar heat equipment

v. Unit heater

Intercom system

Laboratory plumbing, piping

Magnetic door holders

Nurse call system

Oxygen, gas, air piping

Paging system

Plumbing, composite

a. Fixtures

b. Piping

c. Pump

d. Water heater, commercial

e. Water storage tank

Pneumatic tube system

Sprinkler and fire protection system

a. Fire alarm system

b. Fire pump

c. Smoke and heat detectors

d. Sprinkler system

e. Tank and tower

Sewerage, composite

a. Piping

b. Sump pump and sewerage ejector

Telephone system

Television antenna system

Vacuum cleaning system

Water wells

B. Other Fixed Equipment

Bench, bin, cabinet, counter, shelving,

Cabinet, biological safety

Carpentry work

Caulking

Ceramic tile

Conveying system

Drilled piers

Fire protection in hoods

Generator set

Hood, fume

ICU-CCU counters

Lockers, built-in

Mailboxes, built-in

Millwork

Nurses' counter

Painting

Pass-through boxes

Patients' wardrobes and vanities Sink and drainboard

Sterilizer, built-in

X-ray protection

III. Moveable Equipment

Accelerator

Accounting/bookkeeping machine

Acculab

Adding machine

Air conditioner, window

Alternating pressure pad

Ambulance

Amplifier

Analyzer a. Amino acid

b. Autos

c. Biochromatic

d. Clinical

e. Gas

f. Oxygen

g. pH gas h. Peripheral

i. Pulmonary function

Anesthesia unit

Ankle exerciser

Apparatus

a. Anesthesia

b. Blood transfusion

c. Bone surgery

d. Resuscitating

Arthroscopy instrumentation

Aspirator

Audiometer

Autoclave

Automobile

a. Delivery

b. Passenger Autoscaler, ionic

Auto suture stapler

Bassinet, heated Bath a. Paraffin

b. Serological

a. Analytical

b. Electronic

c. Sitz

Balance

Rassinet

d. Water, laboratory

c. Precision mechanical

Basal metabolism unit

e. Whirlpool

Battery charger

Red

a. Electric

b. Flotation therapy

c. Hydraulic

d. Labor

e. Manual

f. Orthopedic

Bedpan washer

Beepers, paging

Bench, metal or wood

Bilirubin lamps

Bin, metal or wood

Binder, punch machine

Biochemical analysis unit, micro

Biofeedback machine

Bipolar coagulator

Blanket drier

Blanket warmer

Blood chemistry analyzer, automated

Blood cell counter

Blood gas analyzer

Blood gas apparatus, volumetrics

Blood warmer

Blood warmer coil

Boiler, copper

Bookcase, metal

Bottle washer

Bovie unit

Breathing unit, positive pressure

Broiler

Bronchoscope a. Flexible

b. Rigid

Buffer, electric

Bulletin board

Burnisher, silverware

Cabinet

a. Bedside

b. File

c. Instrument

d. Metal or wood

e. Pharmacy

f. Solution g. X-ray

Cage, animal

Camera

Camera, surgical

Camera, TV monitoring, color or black

and white

Camera, videotape, color or black and

white

Can opener, electric

Canopy, ventilating, ironer

Capsule machine

Carbon monoxide recorder/detector

Cardioscope

Carpeting

Cart

a. Food/tray, heat-refrig

b. Maid

c. Medicine

d. Supply

e. Utility

Cash register

Cassette changer

Cautery unit

a. Dermatology

b. Gynecology

Central processing unit

Centrifuge

Centrifuge, refrigerated

Chair

a. Dental

b. Executive

c. Hydraulic, surgeon's

d. Kinetron

e. Podiatric

f. Specialist

Chart rack

Chart recorder

Check signer

Child immobilizer Chloridiometer

Chromatograph, gas Cidematic washer

Clock

Clopay wrapping machine Clothes locker

a. Fiberglass or metal b. Laminate or wood

Cobalt unit

Coffee maker

Cold pack unit, floor

Collator, electric

Collector, silver, automatic Colonoscope

Colorimeter Colposcope with floorstand

Compactor, waste

Compresser, air

Computer assisted system for exercise Computer

a. Cardiac output

b. Clinical c. Large

d. Micro

e. Mini

Computer terminal Conductivity tester

Conveyor system, laundry

Conveyor, tray

Cooker, pressure, food

a. Walk-in, freestanding

b. Water

CO-oximeter

Coulter counter

Copier

Credenza Crib

Cryo-ophthalmic unit with probes

Cryostat

Cryosurgical unit Cutter

a. Cloth, electric

b. Food

Cystic fibrosis system

Cystometer Cystoscope

Data card processing unit, including keypunch, verifier, reader, sorter

Data printing unit Data storage unit a. Mechanical

b. Nonmechanical

Data tape processing unit, including controller, drive, tape deck

Decalcifier Defibrillator

Densitometer, recording Dental drill with syringe

Dermatome

Desk, metal or wood

Diagnostic set Diathermy unit Dictating equipment

Digital fluoroscopy unit Digital radiography unit

Dilutor Dish Sterilizer Dishwasher Disinfector

Dispenser a. Alcohol

b. Butter, refrigerated c. Milk or cream

Distilling apparatus

Dopplers Dose calibrator

Dresser Drier

a. Clothes b. Hair c. Sonic Drill press

Drying oven, paint shop

Duplicator

Echocardiograph system

Echoview system Electrocardioscanner Electrocardiograph Electroencephalograph Electromyograph

Electronic blood pressure device

Electrophoresis unit Electrosurgical unit

Emission computer tomography (ECT)

scanner Enlarger Ergometer

Ether-suction unit

Evacuator

Evoked potential unit Exercise apparatus

Extracorporeal shock-wave lithotripters

Extractor, laundry Facsimile transmitter Fiber optic equipment

Fiberometer Film changer Flame photometer Floor scrubbing machine Floor waxing machine

Fluorimeter Fluoroscope Folder, flatwork Food chopper Frame, turning Freezer, ultra cold Fryer, deep fat Furnace, laboratory

Furniture a. Central supply b. Dietary

c. Housekeeping d. ICU-CCU

e. In-service education f. Labor-delivery

g. Laboratory

h. Lobby or public areas

i. Nursing services

Office

j. Office k. Operating room l. Patient m. X-ray Gamma camera Gamma counter Gamma wall system Garbage disposal Graphotype Griddle Grinder, food waste

Hand dynamometer Heart-lung system Hemoglobinometer Hemodialysis unit Hemophotometer Hoist, chain or cable Holter electrocardiograph Holter electroencephalograph

Homogenizer Hood, exhaust or Bacti

Hot-food box Hotplate Humidifier Hyperbaric chamber Hydrocollator

Hydrotherapy equipment

Hyfrecator

Hypothermia apparatus Ice-cream freezer

Ice-cream storage cabinet Ice-cube making equipment Illuminator unit, multifilm Illuminator unit, single

Image intensifier IMI infant care center Immuno-diffusion equipment

Imprinter, addresser Imprinter, embossed plate

Incubator a. Laboratory

b. Nursery Indicator, remote Infusion pump Inhalator

Instruments, ortho-urological

Insufflator Integrator Intercom

Iontophoresis unit Ironer, flatwork Isodensitometer Isolation chamber Isotope equipment Kettle, steam jacketed

Kiln K-pads Kymograph

Laminar air-flow unit

a. Cabinet b. Wall Lamp

a. Deep therapy b. Emergency c. Infrared

d. Mercury quartz e. Microscope

Laparoscope Laryngoscope Lathe

Lawn mower, power Library furnishings Lifter, patient Light

a. Delivery b. Examining c. Operating

d. Portable, emergency

Linear accelerator

Linen cart Linen drier Linen press Linen table Linen washer Loom

Lowerator

Magnetic resonance imaging Mailing machine Mannequin Marking machine

Meat chopper Medi-prep Meter, pH Microfilm unit Microgasometer Microphone Microscope

Microprojector Microtome Mirror, therapy

Mixer, commercial type Model, anatomical

Monitor a. Apnea

b. Cardiac

c. Cerebral function

d. Patient e. TV

Narcotic safe

Natural childbirth backrest

Nebulizer a. Pneumatic

b. Ultrasonic Nephroscope

Neurological surgical table headrest

Nourishment ice station

Nuclear magnetic resonance scanner

Operating stool Opthalmoscope Optical readers Orthotron system Oscilloscope Osmometer Otoscope Ottoman Oven a. Baking

b. Microwave c. Paraffin

d. Roasting e. Sterilizing

Oximeter

Oxygen tank, motor, and truck

Pacemaker, cardiac Pacing system analyzer Packaging machine Paint spray booth Paint spraying machine

Panendoscope Paper baler Parallel bars Parking lot sweeper

Patient monitoring equipment Peeler, vegetable, electric

Percussor Perforator Phonocardiograph Photocoagulator Photocopier

Photography apparatus, gross pathology

Photometer Phototherapy unit

Physicians' in-and-out register, portable

Physiological monitor

Physioscope Piano

Pipe cutter-threader Pipette, automatic Planer and shaper, electric

Plasma freezer Plate bending press Polisher, floor

Polishing and buffing machine

Power supply Press, laundry Proctoscope Projection machine Projector, slide

Prothrombin timer, automated Pulmonary function equipment

Pulsed oxygen chamber Pump

a. Breast b. Stomach c. Surgical

d. Vacuum

Radiation meter Radioactive source, cobalt

Radiographic duplicating printer Radiographic fluoroscopic combination

Radiographic head unit Range, household Ratemeter, dual

Recorder

a. Laboratory b. Tape

Refractometer Refrigerator

a. Blood bank

b. Commercial c. Domestic

d. Undercounter

Remote control receiver

Respirator Resuscitator Retractor Rhinoscope Rinser, sonic Rotary tiller

Roto-osteotome unit

Safe Sanitizer Saw

a. Autopsy b. Band

c. Bench, electric d. Meat cutting

e. Surgical, electric

f. Neurosurgery

Scale

a. Baby b. Bed

c. Chair d. Clinical

e. Laundry, platform

f. Laundry, movable

g. Metabolic h. Patient

i. Postal

Scanner

a. Body CT

b. Isotope c. Rectilinear

d. Ultrasonic Scintillation scaler Sectocardiograph

Sensitometer Seriograph, automatic

Settee

Sewing machine Shaking machine

Sharpener, microtome knife Shears, squaring, floor

Shelving, portable, steel Shoulder wheel

Sigmoidoscope Silver recovery unit

Simulator

Skeleton Slicer

a. Bread b. Meat Slide projector

Slide strainer, laboratory

Slit lamp Snow blower Sofa

Spectroscope Spectrophotometer Sphygmomanometer

Spirometer Stall bars Stamp machine Stand

a. Basin

b. Irrigating

c. IV

d. Mayo

Stapler, electric or air Steam pack equipment Steamer, vegetable

Stencil machine

Stereo equipment Sterilizer, movable

Stethophone Still, water

Station system Stimulator, muscle

Stretcher Suction pump

Table

a. Anesthetic

b. Autopsyc. Electrohydraulic tilt

d. Examining e. Fracture

f. Food preparation

g. Instrument

h. Light i. Metal

i. Obstetrical

k. Operating

1. Orthopedic m. Overbed

n. Pool

o. Refrigerated

p. Therapy q. Traction

r. Urological s. Wood

Tank

a. Cleaning

b. Full body

c. Hot water d. Paraffin e. Therapy

Telemetry unit

Telescope, microlens Telescopic shoulder wheel

Telethermometer Television receiver

Tent

a. Aerosol b. Oxygen

Test equipment

Thermometer, electronic Thyroid testing equipment Time recording equipment

Tissue embedding center Tissue processor

Titrator, automatic Toaster, commercial type Tonometer

Totalap Traction unit Tractor

Transcribing equipment

Transcutaneous nerve simulator system

Treadmill, electric Truck, hot food

Truck a. Forklift

b. Multipurpose filling

c. Van, pickup

Trunk

a. Platform

b. Tray Tube dryer Tube tester

Tumbler

Typewriter

a. Electric b. Manual

Ultrasonic cleaner

Ultrasonic fetal heart detector

Urn, coffee Vacuum cleaner Vacuvette

Valet, office Vectorcardiograph Vending machine

Ventilator, respiratory

Vial filler Vibrator

Victoreens meter Vise, large bench

Walkie-talkie

Warmer

a. Dish b. Food

Washer, glassware Washing machine

a. Commercial

b. Domestic

Water cooler, bottle or fountain type Water purifier or softener

Welder Wheelchair

Wire tightener-twister

Word processor a. Large

b. Small

X-ray

a. Developing tank

b. Film drier

c. Film processor

d. Image intensifier

e. Intensifying screens

f. Wiring

g. Unit, deep therapy

h. Unit, fluoroscopic

i. Unit, mobile

j. Unit, radiographic

k. Unit, superficial therapy

### Appendix D—Regulatory Impact Analysis

# A. Introduction

Executive Order (E.O.) 12291 requires us to prepare and publish a final regulatory impact analysis for any final regulation that meets one of the E.O. criteria for a "major rule"; that is, that will likely result in: An annual effect on the economy of \$100 million or more; a major increase in costs or prices for consumers, individual industries, Federal, State, or local government agencies, or geographic regions; or significant adverse effects on competition, employment, investment,

productivity, innovation, or on the ability of United States-based enterprises to compete with foreign-based enterprises in domestic or export markets.

In addition, we generally prepare a final regulatory flexibility analysis that is consistent with the Regulatory Flexibility Act (RFA) (5 U.S.C. 601 through 612), unless the Secretary certifies that the final regulation will not have a significant economic impact on a substantial number of small entities. For purposes of the RFA, we treat all hospitals as small entities.

We believe that this final rule will result in significant changes in the manner in which hospitals finance their capital expenditures. Accordingly, we have prepared the following discussion, which, in combination with the preamble of this final rule, constitutes a combined regulatory impact analysis and regulatory flexibility analysis meeting the requirements of E.O. 12291 and the RFA. This discussion includes our responses to comments related to the initial analysis published with the May 19 proposed rule.

# B. Summary of the Initial Analysis Published May 19, 1987

In the May 19, 1987 proposed rule, we presented an initial regulatory impact analysis that compared projected payments under the phase-in of prospective payments for capital costs over the applicable transition period (ten years for fixed plant/equipment and two years for moveable equipment) to projected payments under current reasonable cost principles. The comparison used FY 1984 hospital capital cost data that were inflated through FY 1988 by an industry-wide inflation rate developed by the American Hospital Association. We displayed the effects of implementing the proposed payment system over the transition period in terms of the projected percent change in payment levels between the amounts that hospitals would receive under the current system for FY 1988 compared to what they could expect under the proposed capital prospective payment system.

As a result of both the relatively short transition period of two years for moveable equipment and the absence of an exceptions policy in the May 19, 1987 proposed rule, some hospitals might have experienced substantial reductions in their payments for capital costs. Many commenters, in responding to the proposed rule, indicated concern with the proposed payment system. We are now presenting a final regulation that we believe responds to the commenters' concerns.

C. Objectives

The chief objective we hope to achieve through integrating payments for inpatient capital costs into the prospective payment system is to establish the same kind of economic relationship between hospital operational characteristics and market conditions on the one hand, and capital investment decisions, on the other hand, as exists in a price-competitive market. The retrospective payment system now in effect does not constrain hospital capital spending sufficiently to bring these costs under control. Under the present system of capital payments, hospitals may gain access to financial markets for the purpose of obtaining capital and acquiring assets, even if these assets do not contribute to the effective or efficient operation of the facility. For example, under cost-based reimbursement, a hospital with low utilization could borrow funds for expansion of its plant even though it may have surplus capacity, and Medicare would still reimburse those capital costs. This final regulation establishing prospective payment rates for inpatient hospital capital costs, therefore, will establish a payment system that results in hospitals accepting a greater degree of risk for their investment decisions. Payments for capital costs will now face the same financial and economic incentives to which operating costs are now subject.

Comment: Many commenters expressed concern that incorporating reimbursement for capital costs into the prospective payment system might significantly harm their cash flow and overall reimbursement. In particular, several hospital associations asserted that our proposed rates would undercompensate hospitals capital costs and thus would have serious effects on their viability. Others noted that, based on the initial impact analysis, the new payment system may initially benefit some regions or provider types at the expense of others.

Response: We are issuing these final regulations to achieve our objective of more economical capital decisionmaking by health care providers. Moreover, although a number of commenters expressed dissatisfaction with our analysis, only a few commenters presented any detailed data in support of their claims. We are especially disappointed that none of the major hospital associations submitted pertinent national data.

Regarding our impact analysis, we caution readers that data limitations and necessary methodological

assumptions limit our understanding of the effects of these regulations on aggregate groups of hospitals. Even at this level, the results should be interpreted cautiously. For example, some hospitals of a type that appears to suffer reductions in payment in this analysis may not have made any major capital investments between FY 1984 and FY 1988. While our analysis assumes that the inflation-adjusted payments for these hospitals have grown at the same rate as the universe of hospitals between FY 1984 and FY 1988, their actual capital payments may have grown at a slower rate over that period. Ultimately, the requirements of budget neutrality under section 9303(a) of Pub. L. 99-509 require that any increases in payments to some hospitals be offset by payment reductions to others.

Moreover, we believe that the additional payments provided under § 412.68, in conjunction with transition periods for fixed plant/equipment and moveable equipment that are heavily weighted towards the hospital-specific payment in the first few years, should address many of the potential financial problems that may arise during the transition.

# D. Analytical Methodology

### 1. General Considerations

Our approach for displaying the effects of the new payment system is similar to the one we adopted in the May 19, 1987 proposed rule. We will compare estimated FY 1988 payment levels under the new payment system using the FY 1988 Federal payment blends with projected payments under the present system. We are computing projected payments under reasonable cost principles by inflating each hospital's FY 1984 costs through FY 1988, using an estimate of the actual rate of inflation for capital costs. Hospital groupings that will, on average, receive higher payments under the new system will have positive values in our analysis while hospital groupings that will, on average, receive lower payments under the new system will have negative values. These positive and negative values represent the percent increase or reduction in payments compared to projected payments under the present system.

Comment: Several commenters noted that the annual update factors used to inflate the capital expenses of each hospital from FY 1984 to FY 1988 levels were omitted from the proposal.

Response: The updating factors used represent industry-wide estimates of the annual rate of inflation for per-case

inpatient capital costs developed by the American Hospital Association, adjusted for the decline in Medicare inpatient utilization since FY 1984. We have published these estimates elsewhere in this preamble.

Section 1886(g)(3) of the Act (added by section 9303(a) of Pub. L. 99-509) requires payments for capital costs to be reduced by seven percent for portions of cost reporting periods or discharges occurring in FY 1988, regardless of the method of payment. Accordingly, we reduced our projected payments under reasonable cost reimbursement by seven percent. A similar reduction has already been made in the Federal prospective payment rates, so we are maintaining comparability between reasonable cost payments and prospective payments. As a result, the impact analysis in this document assumes that both the Federal payment rates and projected payment rates already have the seven percent removed from them.

Since the hospital-specific portion of the prospective payments for capital is, under these regulations, the hospital's actual allowable costs, we need not compute it separately from the reasonable costs used to simulate payments under the current system. That is, to compute the hospital-specific portion for a given year of the transition. all we need do is multiply the hospital's reasonable costs by the applicable blend factor in effect for that year of the transition. For example, in FY 1988, hospital-specific payments would equal the product of the same capital reasonable costs used to compute reasonable cost payments, multiplied by the applicable blend factor of .95.

In interpreting the results of this comparison, readers should keep in mind the following points:

· The first point is the static nature of the analysis. Hospitals and other interested parties should interpret the following analysis as indicating the direction and magnitude of changes in payment amounts based on the capital costs for hospitals in FY 1984. Although the cyclical nature of hospital investment would result in projected rates of growth in Medicare capital costs per case which are different for each hospital, we did not have the hospital specific data to incorporate these different growth rates in projecting FY 1988 hospital capital costs. Thus, for all of the impact analyses, we increased each hospital's FY 1984 capital costs per case by our projections of the national average increase in Medicare capital costs per case between FY 1984 and FY 1988. While applying this assumption to large aggregates of hospitals allows us

to achieve a reasonable projection of the initial redistributional effects of these regulations, we warn the reader that, in assessing the following tables, the impact results for each group of hospitals assume FY 1984 conditions which may not exist in FY 1988.

• Second, we are presenting the effects of the new hospital payment system for both the FY 1988 Federal blends proposed in the NPRM and for the new blending rate and exceptions policy of this final regulation. In both instances, the comparison is made to our projections of the distribution of capital payments under the current cost reimbursement system, and our estimates of impacts are based on the assumption that the new payment system will take effect on October 1, 1987.

· A positive or negative value associated with a specific hospital grouping does not necessarily mean that hospitals in that group would experience either accounting or economic profits or losses for all inpatient care services. Our results show only decreases or increases of Medicare payments for capital costs under the new payment system, relative to payments under the present system (including the mandated reductions under section 9303(a) of Pub. L. 99-509). At present, Medicare covers, on average, approximately 41 percent of all inpatient hospital costs (based on estimated FY 1986 Medicare data compared to total hospital inpatient revenues as reported by the American Hospital Association), and capital expenditures account for about ten percent of all hospital inpatient expenditures. Thus, a five percent reduction in Medicare capital payments for an average hospital would result in a payment reduction of about .2 percent compared to total inpatient expenses. A hospital could receive lower payments under the new system, but overall, it could still be profitable because of operating gains earned from noncapital services to Medicare patients as well as to other patients. The converse may also occur; hospitals earning surpluses under the capital payment system may be suffering operating losses.

• The final point is that the impact values shown in Table I of Appendix D are single point estimates for different groups of hospitals rather than a range. Since a range of impact values allows one to better understand variations in capital payments among individual hospitals, we are including an analysis of range data for urban and rural hospitals, showing the distribution of hospitals using the blended Federal payment share effective for the first year

of the phase-in (see Table II of Appendix D, which depicts the effects of this final rule in comparison to the

proposed rule).

Comment: One commenter argued that, if the Federal rates are updated for FY 1990 onward using the prospective payment update factor for all other inpatient operating costs, then the new payment system will extend the Pub. L. 99–509 mandatory reductions in aggregate capital spending indefinitely into the future (Section 9303(a) of Pub. L. 99-509 currently does not address national aggregate capital spending levels for FY 1990 and beyond). The commenter alleged that the initial regulatory impact analysis presented with the NPRM "obscures" this regulatory change.

Response: We are presenting our impact analysis of this regulation in terms of its possible FY 1988 distributional effects and other impacts as compared to current law and regulations. Current law provides for a nationwide seven percent reduction in capital-related reimbursement in FY 1988 and a ten percent reduction in FY 1989, regardless of payment mechanism

used.

The data limitations and assumptions inherent in our analytic model, coupled with incomplete information on provider behavioral response to this regulation, forces us to present only short term impact projections here. Subject to these limitations, we nevertheless believe that the analysis contained herein fairly represents the impacts of the new payment system (in and of itself), and that possible Congressional action regarding national aggregate reimbursement levels after FY 1989 remains a separate issue.

Comment: Three commenters noted that the Federal rates for the new payment system are based upon cost and utilization data derived from hospital cost reporting periods that began in FY 1984. Given the fact that the new rates are on a per discharge basis, and given the decline in Medicare inpatient discharges since FY 1983, these commenters asked whether HCFA's impact estimates incorporate any potential decline in revenue that hospitals face under the new payment system on this account.

Response: The rates are based upon our best estimates of total Medicare capital costs in FY 1988. These estimates include the effects of nationwide reductions in hospital occupancy rates that have occurred since FY 1984.

Comment: Commenters expressed concern about possible conflict between HCFA's base year assumptions and those implicit in the proposed transition period for moveable equipment. In particular, one commenter asked

whether the policy and impact analysis will take into account HCFA's guidelines to intermediaries regarding the audit treatment of depreciation expense. Intermediaries evaluate reported capital costs for moveable equipment on the basis of AHA depreciation schedules which assign a longer useful life to most items than was allowed for under the NPRM's proposed transition period.

Response: We believe that extending to seven years the transition period for incorporating capital payments for movable equipment into the new payment system more appropriately accounts for the true depreciable life of these items.

Comment: Several commenters argued that we should present impact projections for blending rates representing the fourth year and beyond of the transition.

Response: We are presenting information giving the direction and magnitude of changes which interested parties can expect to occur in the short term as a result of this rulemaking. However, providers can mitigate the long term impacts of the regulations through several strategies. In the absence of quantified models of provider response to the new payment system for capital, we believe that it would be speculative to present in tabular format impact scenarios for the Federal payment blends for FY 1991 and beyond. Actual adverse impacts can be expected to be much less substantial than shown in the tables, as providers adapt their capital purchasing to the new payment system. Similarly, those hospitals that would be advantaged by the new payment system can be expected to act so as to optimize their benefit.

Comment: Some commenters argued that, in order to perform a full impact analysis, we should provide the projected impacts of the new payment system on a hospital-specific or statewide basis.

Response: As explained earlier, the cyclical nature of capital investments can produce wide fluctuations in capital spending, which we are unable to model, within a short period of time. For an individual hospital, our FY 1988 projection of capital costs could be inaccurate, and an impact analysis for individual hospitals that relied on these data also would be inaccurate. Similarly, we felt that presenting the data on a statewide basis might also be misleading, particularly for States with relatively few hospitals.

The analysis showing the projected range of first year impacts on all hospitals is presented in section D.3.

The analysis in D.3 also presents, for comparison, the estimated first year impacts under our original May 19, 1987 proposed rule.

2. Projected Differences Between Current and New Payment Methods

Table I displays the effects of the new prospective payment system compared to the present system. As previously explained, the results are based on FY 1984 costs inflated to FY 1988 and incorporate both the seven percent reduction mandated under section 1886(g)(3)(A) of the Act and the budget neutrality factor for FY 1988 mandated under section 1886(g)(3)(C) of the Act. It should be noted that, since we have no data on the investment patterns for each hospital between FY 1984 and FY 1988, the impact values for various groups may not be representative of the actual gains or losses resulting from our prospective payment system.

Comment: Many commenters expressed concern regarding the impact of the proposed short transition to a 100 percent Federal rate for movable

equipment.

Two commenters asserted that Table I should present the impacts for the fixed plant/equipment and movable equipment provisions of the proposal separately. Another suggested that HCFA assess the impact of the split transition period for fixed plant/ equipment and movable equipment on hospitals with different relative investments in the two categories.

Response: The transition period for movable equipment has been lengthened considerably to seven years in this final rule, and for the first four years the Federal blends for both fixed plant/ equipment and movable equipment are the same. Accordingly, we believe that most commenters' concerns regarding the split between fixed plant/equipment and movable equipment have been addressed.

We cannot present with any degree of confidence the impacts of the several provisions of this rule separately, since many hospitals probably have different relative investments in the two capital categories in FY 1988 than they had in FY 1984. For this reason, we believe it most appropriate to present our projections of the impact of this rule on hospitals' total capital payments, especially given the fact that the Federal blend for the two capital categories will be equivalent in the first four years. We originally presented the impact of the proposed rule on hospitals' total capital expenditures because we believe that many hospitals are most concerned about the impact of this regulation on their total reimbursement levels.

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MOVEABLE CAPITAL COST IN FY 1988 COMPARED  TO ESTIMATED IMPACTS OF THE FIRST YEAR BLEND AS PROPOSED MAY 19, 1987 1/  Fixed - 5% Federal  Moveable - 33% Federal  OF Moveable - 5% Federal  OF (PROPOSED PAYMENT RATES) 3/  (FINAL PAYMENT RATES)	0.0		2.6		1.9				2.0		4	4:		9	.2	0.0		6.	.3	-0.2	-0.6	0.5	Story and I have a second or the second
MOVEABLE C TO ESTIMATE AS AS Number of Hospitals 2/ (P	All Hospitals 4820	Urban by Region	total side	South Atlantic 362 Fast North Central 483		Central 188	461	Rural by Region	30	Mid Atlantic 80	tral		Central 404	89	Urban Hospitals 2554		1501	tals 2212		Beds 347	Teaching Status 3824 Non-Teaching	Resident/Bed Ratio	0

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2	
18	
6	7
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continued

1

Table I

(PROPOSED PAYMENT RATES) Moveable - 33% Federal 5% Federal

Fixed - 5% Federal Moveable - 5% Federal (FINAL PAYMENT RATES)

Hospitals 2/ Number of

Disproportionate Share Hospitals (DSH)
No Additional Payments

0.5

1.4

82

793

Urban DSH fewer than 100

Rural DSH

Beds

Beds or More Urban DSH 100

00

0

0 0

196

Other Special Status Rural Referral Centers

Rural fewer than

50 beds (RRCs)

0.5

3.3

712

Type of Ownership Voluntary

Proprietary Government

This table represents estimated Medicare margins for the Federal/hospital-specific payment blend for fixed and moveable capital costs for the first year of the transition from the present capital payment system to the prospective payment system compared to Medicare margins that were estimated based on the prospective payment rates for fixed and movable equipment proposed in the May 19, 1987 projected payments under the new payment system to payments under the current system, based on 1984 proposed rule as corrected by the June 11, 1987 notice. These margins result from a comparison of data inflated through FY 1988, and assumes all hospitals have cost reporting periods corresponding to the beginning and end dates of the phase-in year. Both sets of margins reflect a budget

neutrality factor based on projected FY 1988 payments and the exclusion of sole community hospitals from the proposed payment system.

2/

This column shows the number of hospitals in each cell of the FY 1984 hospital cost data base used to compute both the proposed payment rates and the estimated impact. Puerto Rican hospitals are only included in the national hospital total of 4820; otherwise, Puerto Rican and sole community hospitals are not included in these figures. Margins based on a blend of 95 percent hospital-specific for plant/fixed equipment, and 67 percent hospital-specific portion for the moveable equipment that was proposed in the May 19, NPRM, as updated by revisions to the notice published on June 11, 1987. 3/

Margins based on a blend of 95 percent hospital-specific portion for both plant/fixed equipment. and moveable equipment. 4

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Table I clearly demonstrates that the first-year impacts of this final rule will be much less severe than the projected effects of implementing the May 19, 1987 proposed rule without change. For every region and every hospital category shown in our analysis, the aggregate first year capital payments which hospitals will receive under the new payment policy will more closely approximate their projected payments under cost-based reimbursement, than would have been the case under our proposed rule. We attribute this change to the lengthened transition period for moveable equipment and the inclusion of the exceptions policy for hospitals with high capital costs

Regarding the specific impact of our capital payments exceptions policy, we project that slightly over 13 percent of all hospitals will qualify for these payments. We project these payments in the first year to total about 10 million dollars nationally, and these exceptions payments will increase during the transition period as the Federal payment blend increases. Accordingly, our projections show that about 16 percent of urban hospitals and almost 11 percent of rural hospitals will qualify for additional payments.

In the aggregate, our analysis of this final rule as presented in Table I shows no change in overall payments from the present method of paying for inpatient hospital capital costs. This is to be expected because we have maintained throughout this analysis the budget neutrality requirement imposed on payments for the first year. Nevertheless, the impact varies among geographic regions Based on the first year Federal blend the impact ranges from a high of 2.0 percent increase for rural hospitals in the New England region to a low of 0.8 percent decrease for urban hospitals in the Mountain census division.

Among categories of hospitals there is also a wide range of projected impacts. Rural hospitals, overall, fare better than urban hospitals, with rural hospitals that have fewer than 50 beds doing especially well, showing, on average, a 1.3 percent increase in payments. All rural hospitals, in the aggregate, show a 0.1 percent average increase as compared to an insignificant decrease (less than 0.05 percent) in the aggregate payments for urban hospitals. It appears that the gains made by small rural hospitals are, in part, attributable to our computing the standardized payment rates on a case-weighted basis in accordance with section 1886(d)(3)(A) of the Act (as amended by section 9302(c) of Pub. L. 99-509).

Similarly, teaching hospitals with "heavy" graduate teaching programs (defined as hospitals with resident to bed ratios of 0.25 or greater) show an average increase in their capital payments of approximately 1.1 percent when projected payments are based on the first year Federal payment blend. Other categories of hospitals that would receive higher payments under the proposed system are large urban hospitals (with 405 beds or more) and government-controlled facilities. The latter group is shown to benefit by a 1.1 percent increase based on the first year blend.

Among hospitals that are likely to experience a drop in payments are proprietary facilities, with a projected payment decrease, under the first year Federal payment blend, of 1.1 percent. The losses resulting from high capital costs for these hospitals will be mitigated by our exceptions policy.

In general, any observed reductions in FY 1988 payments do not necessarily imply losses for hospitals. A noncash expense, such as depreciation, may cause a hospital to show an accounting loss, but does not affect cash flow. Also, principal payments, which Medicare does not reimburse, would not significantly affect cash flow, since these are usually low during the initial years after investment. Finally, hospitals that incur a loss in capital payments in FY 1988 due to a major recent investment will have fairly level capital expenses over the next ten years. Since the capital payments for all hospitals will be increased each year by the prospective payment system update factor, we would expect that a number of these hospitals will obtain capital surpluses during the transition.

Comment: Several commenters disagreed with our assertion that hospitals with initial losses under the new payment system could take action to adapt their capital spending, such as refinancing their debt at lower interest rates. They also questioned our assertion that, as the capital prospective payment rates are updated over time, hospitals which initially receive lower payments would eventually receive adequate reimbursement.

Response: We agree that interest rates appear to be rising at this time. However, we continue to believe that hospitals have the ability to undertake various term financing arrangements that could bring their stream of debt payments over time into line with their expected Medicare capital reimbursement. As stated, most hospitals can choose to postpone some capital expenditures until they have

built an appropriate cash reserve. Also, hospitals could arrange to make heavier debt payments earlier or later in the capital cycle as necessary. We believe that the gradual transition period incorporated in this final rule should allow most hospitals ample time to adjust their capital expenditures to the new payment environment.

We disagree with those commenters who questioned our assertion that hospitals initially receiving inadequate payments will eventually receive adequate reimbursement. If a hospital's debt service remains constant or declines over time (because it is postponing the acquisition of noncritical assets), and our prospective rates are increased over time, then Medicare capital-related payments to such a hospital will eventually more than cover the Medicare share of the cost of related expenditures.

Comment: One commenter argued that the impact analysis given in the proposed rule should examine the interactive effects of the new payment system with Medicaid regulations, in particular those requiring State Medicaid programs to set payments at levels that, on average, do not exceed Medicare levels as an upper bound (§ 447.253(b)(2)).

Response: Under § 447.253(b), Federal financial participation to State Medicaid agencies is limited to an aggregate upper limit for payments to providers equal to the amount Medicare would pay Because section 1886(g)(3) of the Act requires that capital prospective payments be budget neutral with respect to payments under the present system for FYs 1988 and 1989, aggregate Medicare payments to hospitals in some States will decline slightly below present payment levels. Because payment systems vary from State to State (even among those States that generally follow Medicare payment principles), and because States are subject to an aggregate upper limit that covers a number of different types of providers, States have considerable flexibility in responding to the new Medicare prospective payment system. As a result of such flexibility, and the relatively slight changes required by this regulation, we cannot determine how State agencies will respond. Consequently, we cannot determine the fiscal effect of the new systems on Medicaid providers. We believe the most significant payment adjustments State agencies need consider are the seven and ten percent reductions in payments for FYs 1988 and 1989, respectively, mandated under section 1886(g)(3)(A) of the Act.

For periods beyond FY 1989, it becomes increasingly difficult to predict the effects of this rule on Medicare providers. It is well beyond our present analytical capabilities to determine the possible effects of the new payment system on FFP levels to States and how State agencies will respond to changes in such levels.

It is also unclear how the new Medicare prospective payment system for capital costs will affect the amendment of Medicaid State plans.

Following the implementation of the Medicare prospective payment system for inpatient services, many States adopted similar DRG-based prospective payment systems or significantly modified their cost reimbursement systems to incorporate prospective payment features. Significantly, many of the new systems are comprehensive payment systems that include prospective payment for capital costs as well as payments for inpatient operating services. Thus it seems unlikely that the Medicare prospective payment system

for capital costs will stimulate major changes in many Medicaid State plans.

Comment: Several commenters argued that further analyses of the characteristics shared by hospitals that will be adversely affected by the regulations need to be conducted prior to final implementation. In particular, one commenter argued that, prior to the final rule, we should assess whether urban and rural hospitals as groups are in different stages of the capital cycle.

Response: The commenters are, in essence, suggesting that we do an analysis based on the age of hospitals' plant and equipment and capital financing to definitively determine the impact of this rule on hospitals. Since the information necessary to perform such an analysis is not available, it cannot be done. Furthermore, it should be noted that in a competitive marketplace, the price of an item does not vary based on the age of the seller's capital assets. For similar reasons, we believe that prospective payments made to hospitals should not vary on the basis of their age.

3. Projected Distributional Effects of This Regulation

Recognizing that the above table and analysis provide only a measure of central tendency, and fail to give an indication of the range of possible effects, we have examined the range of possible effects the new payment system may have on all hospitals subject to the prospective payment system, and hospitals grouped by their location in either urban or rural areas for the first year of the transition. In presenting this analysis, we must point out again that our analysis does not reflect any investment changes for hospitals since FY 1984. Table II displays both the absolute number of hospitals in our data base groups and the percent they comprise of the number of hospitals in the group under analysis. Again, for the sake of comparison, we also present the range of impacts of the May 19, 1987 proposed rule (as those figures were updated by the June 10, 1987 correction notice published in the Federal Register).

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# TABLE II DISTRIBUTION OF HOSPITALS

Percentage Increases or Decreases in Payments Based on a Comparison of the Proposed Corrected Rates and Final Prospective Payment Rates to Projected Cost-Based Payments 11/

First Year Payment Blend

			Percent	Percent Decreases2/	TV B B B	180	di	Per	Percent Increases2/	ses2/	T 0 12 12 12 12 12 12 12 12 12 12 12 12 12
	Hospitals in data base5/	Less than 20 percent	Less than 20 percent -20 to -15	-15 to -10	-15 to -10 -10 to -5 -5 to 0	-5 to 0	0 to +5	+5 to +10	+10 to +15	0 to +5 +5 to +10 +10 to +15 +15 to +20	Greater than 20 percent
					DISTRIBUTION	DISTRIBUTION OF HOSPITALS BASED ON FINAL RATES 3/	BASED ON FIN	IAL RATES 3/			
All Hospitals	4766	0.0	0.0	0.0	0.0	39.6	40.9	13.7	3.4	1.3	1.1
Urban Hospitals	2554	0.0	0.0	0.0	0.0	46.8	38.2	10.3	2.5	1.3	6.0
Rural Hospitals	2212	0.0	0.0	0.0	0.0	31.2	44.0	17.71	4.5	1.4	1.2
				per Ille	ISTRIBUTION	DISTRIBUTION OF HOSPITALS BASED ON PROPOSED RATES 4/	ASED ON PROP	OSED RATES	bause Mary a pare b		
All Hospitals	4718	9.0	1.3	3.3	10.9	20.9	17.9	14.1	9.5	4.9	15.5
Urban Hospitals	2535	0.5	1.2	3.9	13.2	24.7	19.4	12.7	7.9	5.3	11.1
Rural Hospitals	2183	0.7	1.3	2.6	8.2	16.4	16.0	15.8	10.8	1.1	20.5

- Analysis based on FY 1984 hospital data inflated by an estimate of the actual rate of inflation used in calculating both a projection of payments under current reasonable cost principles and hospital-specific payments under the final (and NPRM) capital payment system. 17
- Increases and decreases are relative to 1984 payments projected to 1988 levels under current reimbursement principles. The rates used for comparison purposes assume that all hospitals in the data base have an October 1 to September 30 cost reporting period. 17

Final payments used in this analysis are based on FY 1988 blend of five percent Federal blend for both plant/fixed equipment and moveable equipment to

simulate payments during the first transition year.

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- Proposed payments used in this analysis are based on FY 1988 proposed blend of five percent Federal for plant/fixed equipment and 33 percent Federal blend for moveable equipment to simulate payments during the first transition year published in the May 19, 1987 proposed rule as corrected in the June 11, 1987 notice. 4
- 5/ Excludes Puerto Rican and sole community hospitals.

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Again, we can see from Table II how the policy changes incorporated in this final rule will mitigate the most serious adverse consequences of our proposed rule. Whereas we project that nearly 16 percent of hospitals nationally would have suffered a first-year reduction in payments of five percent or more under the proposed rule, no hospitals will lose more than five percent in payments compared to cost-based reimbursement as a result of this final rule. Indeed, our final policy prevents such a result, since the first year payment blend incorporates a ninety-five percent hospital-specific amount. As the transition proceeds, the capital exceptions policy and the longer transition period for movable equipment will mitigate the financial impact of the new capital payment system on high capital cost hospitals.

In our analysis of the effects of this final rule, we find that at the national level, for the first year of the proposed system approximately 80 percent of all hospitals in the data base would either lose or gain between zero and five percent compared to current payments for inpatient capital costs. The breakdown between those hospitals that would lose between zero and five percent is almost identical to the percent of hospitals that would gain between

zero and five percent.

A similar clustering around the -5 percent and +5 percent interval is also evident among hospitals grouped by urban and rural locations.

Approximately 75 percent of all rural hospitals fall within this interval, while 85 percent of all urban hospitals will

either lose or gain between zero and five percent. Of all rural hospitals in the data base, nearly 44 percent are projected to receive between zero and five percent higher payments under the proposed payment system for the first year compared to their current payment levels while about 31 percent of rural hospitals would stand to lose between zero and five percent of their current payments. Similarly, among urban hospitals, approximately 38 percent of the hospitals are projected to receive increases of between zero and five percent while about 47 percent of urban hospitals are expected to receive between zero and five percent lower payments.

During FY 1988, about 25 percent of all rural hospitals would receive increases in their payments for capital related expenses of five percent or more, while no rural hospitals will receive payment decreases of 5 percent or greater. About 15 percent of urban hospitals would get payments increases of five percent or

greater.

On the whole, based on policies and payment rates that will take effect in FY 1988, more hospitals would receive increases in payments over current levels than hospitals receiving decreases in their payments. Nationally, we project that about 61 percent of all hospitals in the data base (70 percent of rural hospitals and 53 percent of all urban hospitals) will receive increases in their capital related payments in the first year. Furthermore, almost 20 percent of hospitals will receive payment increases during the first year of more than 5 percent.

Comment: One commenter asked whether the ranges of impacts given in Table II could be given in terms of dollars-per-discharge as well as in percentages. The commenter asserted that this data should be presented for each of the first three years and the final year of the transition.

Response: Generally, we believe it is inappropriate to present in tabular format the range of possible impacts which may occur upon implementation of this regulation for many years into the future. All projected range impacts from the data are particularly sensitive to assumptions regarding changes in hospitals' debt structure between FY 1984 and FY 1988. Also, the addition of an exceptions policy, as well as possible behavioral changes on the part of providers in response to this regulation, will affect the distributional impacts of this policy. However, we cannot at this time model the magnitude of these factors precisely.

We are choosing to give the impacts for Table II in terms of percentages because hospitals can relate these figures to their own particular circumstances. The impact that a given dollar-per-discharge gain (or loss) will have on a particular provider depends in part upon what its average dollar-per-discharge capital costs actually are. For this reason, we believe that presenting this data in terms of dollars-per-discharge gained (or lost) would be less illuminating for most hospitals than percentages.

[FR Doc. 87-20081 Filed 8-28-87; 8:45 am] BILLING CODE 4120-03-M



Tuesday September 1, 1987



# Department of Health and Human Services

**Public Health Service** 

42 CFR Part 59
Statutory Prohibition on Use of
Appropriated Funds in Programs Where
Abortion Is a Method of Family Planning;
Standard of Compliance for Family
Planning Services Projects; Proposed
Rules



### DEPARTMENT OF HEALTH AND HUMAN SERVICES

**Public Health Service** 

42 CFR Part 59

Statutory Prohibition on Use of Appropriated Funds in Programs Where Abortion Is a Method of Family Planning; Standard of Compliance for Family Planning Services Projects

AGENCY: Public Health Service, DHHS.
ACTION: Proposed rules.

SUMMARY: The Public Health Service (PHS) proposes to amend the regulations governing the use of funds for family planning services under Title X of the Public Health Service Act in order to set specific standards for compliance with the statutory requirement that none of the funds appropriated under Title X may be used in programs where abortion is a method of family planning. This change is being proposed to bring the compliance requirements for programs using Title X funds into conformity with the statutory ban on such use of Title X appropriated funds. The proposed amendments should improve compliance by grantees with the statute and facilitate monitoring of compliance by PHS.

DATE: Comments must be in writing and be received by November 2, 1987. It is intended that final regulations will be promulgated within 45 days following the close of the above noted comment period.

ADDRESS: Comments should be sent to the Deputy Assistant Secretary for Population Affairs, Department of Health and Human Services, P.O. Box 23993, L'Enfant Plaza, Washington, DC 20026–3993.

FOR FURTHER INFORMATION CONTACT: Nabers Cabaniss at 202-245-0152.

SUPPLEMENTARY INFORMATION: On July 30, 1987, President Reagan announced that the Department of Health and Human Services would, within 30 days, publish draft regulations governing grants under Title X of the Public Health Service Act, 42 U.S.C. 300, et seq., to give effect to the statutory prohibition on the use of Title X appropriated funds in programs that include abortion as a method of family planning. Set out below are the Department's proposed regulations, along with a statement of the basis and purpose of the amendments. The regulations proposed herein, when they become final, will automatically supersede the present Title X guidelines to the extent those guidelines are inconsistent with the final rules. After the final rules are issued, the Department intends to issue revised Title X guidelines in conformity therewith.

# Background

Title X of the Public Health Service Act was enacted in 1970 by Pub. L. 91-572. Title X authorizes the Secretary of Health and Human Services to, among other things, make grants to public and private nonprofit entities "to assist in the establishment and operation of voluntary family planning projects which shall offer a broad range of acceptable and effective family planning methods and services * * * ." Section 1001(a) of the Public Health Service Act, 42 U.S.C. 300(a). Approximately 95% of the funds appropriated for Title X since enactment have been used to fund family planning service projects under section 1001(a). At present, 90 services grants are funded under section 1001(a); these grants fund the provision of voluntary family planning services at approximately 3,900 clinic sites.

Since enactment, Title X has contained the following prohibition at section 1008:

[n]one of the funds appropriated under this title shall be used in programs where abortion is a method of family planning.

The legislative history of Title X in general, and of section 1008 in particular, reflects a fundamental dichotomy between the provision of preventive and other pre-pregnancy family planning services, on the one hand, and abortion on the other. As was stated in the Conference Report:

[i]t is, and has been, the intent of both Houses that funds authorized under this legislation be used only to support preventive family planning services, population research, infertility services and other related medical, informational, and educational activities. The conferees have adopted the language contained in section 1008, which prohibits the use of such funds for abortion, in order to make clear this intent. Conf. Rep. No. 91–1667, 91st Cong., 2nd Sess. 8–9 (1970).

While the Conference Report reflects the conferees' understanding that certain "medical, informational and educational activities" are authorized under Title X, it is clear that these activities must be "related" to "preventive family planning services, population research, and infertility services." Id. Actions that promote abortion are manifestly distinct from these activities. This distinction is emphasized by the explicit contrast between abortion and family planning drawn in the floor statement of Representative Dingell, the sponsor of section 1008, who stated:

In explaining the purpose of section 1008, Representative Dingell indicated in his floor statements that this provision was intended to prohibit more than the actual conduct of abortions, Rather—

[w]ith the "prohibition of abortion" amendment—Title X, section 1008—the committee members clearly intend that abortion is not to be encouraged or promoted in any way through this legislation. Programs which include abortion as a method of family planning are not eligible for funds allocated through this act. 116 Cong. Rec. 37375 (1970).

He also observed that-

[i]f there is any direct relationship between family planning and abortion, it would be this, that properly operated family planning programs should reduce the incidence of abortion. *Id.*¹

Thus, it is clear that Title X is meant to fund the provision of preventive and other pre-pregnancy family planning services, and not to promote or encourage abortion in any way.

HHS's interpretation of these policies over the years, however, has not provided clear standards for grantees and HHS personnel. In 1982, the Department's Office of the Inspector General (OIG), after auditing 32 Title X clinics, found that the Department's failure to provide specific program guidance regarding the scope of section 1008 had created confusion about precisely what activities were proscribed by the section, and had resulted in variations in practice by grantees. In particular, the OIG audits found that the clinics were relying upon the Department's policy of permitting both Title X family planning services and separately funded abortion-related activities to be provided at a single site. Similar findings were noted by the General Accounting Office (GAO) in an audit of 14 Title X clinics, also conducted in 1982. GAO went on to recommend that "the Secretary establish clear operational guidance by incorporating into the Title X program regulations and guidelines, HHS' position on the scope of the abortion

¹ Regulations implementing section 1008 were initially issued in 1971 (36 FR 18465, Sept. 15, 1971) and revised in 1980 (45 FR 37436, June 3, 1980). In both cases, the regulations stated that Title X projects could not provide abortion as a method of family planning.

restriction in section 1008." 2 The President's July 30th directive to the Secretary was based in part upon this demonstrated need for changes in existing HHS program guidelines.

Accordingly, pursuant to its rulemaking power under 42 U.S.C. 300a-4(a). HHS has herein proposed to revise the regulations governing Title X so as to conform the obligations of grantees to the statutory prohibition in section 1008, and to establish standards for compliance with section 1008 that will permit adequate monitoring of such compliance. Providing Title X grantees with clear notice of specific compliance standards that give effect to the prohibition in section 1008 is necessary to improve compliance by grantees with section 1008 over the level that presently exists. At the same time, issuance of the proposed rules will strengthen the Department's ability to monitor compliance with section 1008 by providing a clearer basis for measuring grantee activities against objective requirements. Finally, the proposed rules will enable HHS to better enforce section 1008 by providing a basis in regulation for disallowance of costs or termination of program funding where noncompliance exists.

# Provisions of the Proposed Rules

Title X authorizes grants for family planning programs. It further specifies that Title X funds may not be used in programs that include abortion as a method of family planning. The rules proposed below set out specific requirements intended to enforce this congressional mandate by making it clear that a Title X program is limited to providing family planning services, and may not provide abortion counseling and referral; that a Title X program must be entirely separate and distinct. financially and physically, from any abortion-related activities; and that a Title X program may not encourage, promote or advocate abortion as a method of family planning.

The proposed regulations, however, apply only to a Title X-funded "program" or "project": that is, "the identified activity approved by the granting agency for support." HHS Grants Administration Manual, Ch. 1-85-20. The proposed rules in no way purport to restrict an organization's activities in programs that are supported otherwise than by Title X funds. Definitions clarifying this terminology are included in the proposed rules. This

program.

Certain provisions of the proposed rules derive directly from and strengthen the Department's longstanding practice in implementing section 1008. See, e.g., proposed § 59.9 (relating in part to financial and accounting separation of abortion-related services from family planning programs) and proposed § 59.10 (relating in part to activities that encourage, promote or advocate abortion as a method of family planning by using legal action to make abortion available as a method of family planning, or by developing or distributing materials advocating abortion as a method of family planning). Since these rules represent no substantial change from prior practice, it would be most helpful if any comments in these areas contain suggestions for improvement based on prior operational experience with existing requirements.

Proposed § 59.7, requiring that programs seeking Title X funding provide an assurance that they will not include abortion as a method of family planning, is procedurally similar to a requirement that appeared in the Title X regulations until 1980. See 42 CFR 59.5(a)(9), as in effect from 1971 through 1980. New § 59.7, however, more closely tracks the language of section 1008, and incorporates more specific requirements designed in part to enable the Secretary to obtain at the application stage information relevant to determining whether a program will in fact comply with the statutory prohibition. If an applicant for Title X funds cannot demonstrate that it will comply with the statutory prohibition by conducting its family planning program consistent with the requirements of each of the proposed rules, it will not be eligible for Title X funds.

Proposed § 59.8 prohibits Title X projects from providing counseling and referrals for abortion. In the past, "mere referral" for abortion and nondirective counseling regarding abortion were not prohibited by the guidelines, on the theory that such activities do not promote or encourage the performance of abortion. Thus, the current Title X program guidelines require that whenever counseling for dealing with unintended pregnancy is requested, family planning clinics funded under Title X must provide "mere referral" and "nondirective" counseling on all options, including abortion, for dealing with the unintended pregnancy. As clearly contemplated by Title X and its legislative history, however, "family planning" is meant to address plans and methods for facilitating or preventing pregnancy, not for terminating it. As such, medical services or counseling related to pregnancy care after pregnancy diagnosis, or any services relating to abortion as a method of family planning, are outside the scope of activities supported by Title X funds.

Moreover, it is clear that counseling activity and other forms of information distribution were understood by Congress to be a significant part of the "family planning services" that Title X funds were to be used to fund. Thus, "mere referral" and counseling are clearly covered by the prohibition in section 1008 on abortion as a method of

family planning.

In addition, it is unreasonable to assume that counseling and referrals for abortion do not indeed "encourage or promote" abortion. Specifically, the purpose of counseling programs for pregnant women is to provide information upon which they will base a course of action; counseling concerning abortion would be pointless in the absence of an expectation that some women receiving such counseling will choose to have an abortion. Similarly, providing a referral for abortion facilitates the obtaining of abortion, and the intended and actual effect of a referral program is that at least some women referred will select abortion as a method of family planning. Thus, even if abortion counseling and referral were not prohibited by the express language of section 1008 as family planning services that include abortion, the statutory purpose of section 1008 not to promote or encourage abortion would be better effectuated by proposed § 59.8.

In order, therefore, to conform program policies with the general statutory limitation on the use of Title X funds for "family planning services" and the specific prohibition in section 1008 on the use of Title X funds in programs where abortion is a method of family planning, as well as to better effectuate

limitation on the scope of the proposed rules reflects the express application of the section 1008 prohibition to "programs," and the statute's legislative history to the same effect. It is also consistent with existing case law holding that the government may favor normal childbirth by refusing to fund or promote abortion, but it may not preclude organizations whose programs receive government funds from using nongovernment resources in other programs that may include abortionrelated activities. The proposed regulations accordingly are not to be construed as restricting or limiting the activities of grantee organizations when such activities are entirely outside of. and separate from, a Title X-funded

² Comp. Gen. Rep. No. GAO/HRD-82-106. "Restrictions on Abortion and Lobbying Activities in Family Planning Programs Need Clarification." p. 22 (1982) (hereafter referred to as the GAO Report).

the statutory purpose of not promoting or encouraging abortion, proposed § 59.8 prohibits abortion counseling and referral, as well as medical services or counseling related to pregnancy care after pregnancy is diagnosed. One of the effects of these regulations will be to insure the ability of otherwise eligible organizations or programs that refuse to engage in abortion-related activities to receive support under Title X.

Although proposed § 59.8 below prohibits counseling or referral for abortion, as well as counseling and other services relating to pregnancy that are provided after pregnancy diagnosis, it should be noted that the current Title X regulations provide, at § 59.5(b)(1), for "necessary referral to other medical facilities when medically indicated." Referrals to a comprehensive list of health-care providers who provide prenatal care and delivery are therefore permitted, provided that such referrals are not used as an indirect means to encourage or promote abortion. However-notwithstanding the Department's past view of this provision as requiring referrals for abortion in cases where it is medically indicated, such as where continuation of the pregnancy would endanger the life of the mother-it is the express purpose of the specific rule changes proposed below to insure that Title X-funded family planning project do not provide counseling or other services relating to the issue of "medical indication" for abortion. Rather, consistent with the legislative intent expressed in Title Xthat is, the provision of preventive and other pre-pregnancy family planning services-\$ 59.8 requires that pregnant women be referred outside of the Title X project for prenatal care and other related medical services. In no case, therefore, should a Title X-funded family planning program make a determination of the appropriateness of abortion.

Read together with proposed § 59.8, § 59.5(b)(1) will thus require referral in any case where pregnancy is diagnosed. Specifically, when a woman who is already pregnant comes to a Title X-funded family planning program, the program must provide her with a full listing of licensed health care providers of appropriate prenatal medical care and delivery services, from which she may select. This requirement is consistent with the legislative design of Title X as a program limited to funding preventive and other pre-pregnancy family planning services.

The Department solicits comments relating not only to proposed § 59.8, but also to its intended effect upon the meaning of § 59.5(b)(1). If necessary, the

Department may amend the language in § 59.5(b)(1) in the final rules, in order to insure that the proposed change is unambiguous.

Proposed § 59.9 articulates new requirements designed to strengthen the Department's existing policy that abortion-related services must be "separate and distinct" from a Title X-funded program. Among these new requirements are provisions relating to the maintenance of separate medical records systems and the physical separation of a Title X project from any abortion-related functions or facilities.

The requirement of proposed § 59.9 that grantees maintain project medical record systems separate from any abortion-related operations is based squarely on the congressional intent that abortion not be a part of a Title X funded program. In this regard, the Department is concerned that commingled data systems may cause grantee organizations to aggregate abortion clients with Title X clients, and may inhibit monitoring of the segregation required by section 1008. The proposed rule thus seeks to ensure clearer records for purposes of excluding abortion-related activities from Title X funded programs and facilitating program monitoring. In fact, there is evidence that this requirement reflects the current practice of some grantees. The Department does not, therefore, anticipate that overall this requirement will impose substantial additional administrative burdens on grantees. See the GAO Report, p. 8.

The provisions of proposed § 59.9 relating to physical separation of abortion activities and family planning programs, while new, effectuate the underlying policy of section 1008. In the past, HHS has not consistently interpreted the statute so as to prohibit situations where the Title X project shares physical facilities (such as a common waiting or treatment area) with a project providing abortion services. HHS has now concluded, however, that a requirement of physical separation is necessary to strengthen the enforcement of the prohibition in section 1008.

In practice, an impermissible use of Title X funds may occur when the physical facility of a grantee organization's Title X-funded family planning program overlaps that of its abortion-related operations. Even where the strictest accounting and charging of expenses is performed, shared facilities inevitably increase the likelihood that a violation will occur, and lead to situations where the assertion that a program does not "include" abortion

amounts to little more than an accounting fiction.

Accordingly, one purpose of proposed § 59.9 is to insure that Title X funds not be used for abortion-related activities. In addition, it is intended to further enforcement of the statutory requirement of section 1008 that abortion not be a method of family planning in a Title X program. Meeting this latter requirement mandates that Title X programs be organized so that they have an appropriate integrity and independence from other activities conducted by the grantee which are prohibited by statute from inclusion in a Title X funded program. Having a program that is separate and distinct from other such activities conducted by the grantee is a necessary predicate to any determination that abortion is not being included as a method of family planning in the program.

Moreover, proposed § 59.9 is independently justified by the need to prevent existing or potential clients of Title X projects—as well as the general public-from concluding that the government endorses abortion. By promoting the view that abortion is an acceptable and government-sanctioned method of family planning, moreover, the rendering of abortion and family planning services in common facilities violates the intent of Congress underlying section 1008, that Title X funds will not be used to "encourage or promote" abortion. Thus, proposed § 59.9 prohibits siting a Title Xsupported family planning program in a fashion which would result in use of shared physical facilities-for example, with respect to waiting, consultation. examination, and treatment areas. It also prohibits Title X-funded projects from sharing office entrances and exits with an abortion facility. These proposed requirements effectuate the policy expressed in section 1008 that Title X projects not include abortion as a method of family planning.

One additional provision belowproposed § 59.10(a)(1) (relating to payment of dues to advocacy organizations)—constitutes a change from current program requirements. The provision of proposed § 59.10(a)(1) prohibiting payment of dues with project funds to advocacy organizations is necessary to ensure that Title X funds are not indirectly used to advance objectives that are not only inconsistent with Title X, but specifically prohibited by section 1008. Absent the restriction in proposed § 59.10(a)(1), neither the Department nor the grantee could ensure that Title X funds will not be used to encourage or promote

abortion—activities which are prohibited by section 1008. See the GAO Report, pp. 24–26.

# Regulatory Flexibility Act and Executive Order 12291

The proposed rules codify existing statutory requirements applicable to Title X grantees. With one exception. the effect of the proposed rules is to eliminate existing requirements or permissive provisions concerning the provision of abortion-related services. and as a result the proposed rules should to this extent produce a reduction in costs for Title X-funded programs. The exception is proposed § 59.9, relating to separation of abortionrelated services from family planning programs. According to the Department's information, approximately 80 of the approximately 3,900 Title X-supported family planning sites are physically located near facilities that provide abortion services. Of these 80, it is unknown how many currently meet the requirements of proposed § 59.9. However, in view of the fact that the potential number of sites affected is small, and in view of the fact that current requirements under Title X already prohibit any direct subsidy of abortion services with Title X family planning funds, the Department believes it is unlikely that the proposed rule would have economic consequences even approaching the threshold for major economic consequences as defined in Executive Order 12291.

For the foregoing reasons, and consistent with the provisions of the Regulatory Flexibility Act [5 U.S.C. 605(b)], the Secretary also certifies that this rule will not have significant economic impact on a substantial number of small entities.

# Paperwork Reduction Act

Proposed § 59.7 and proposed § 59.9 contain collection of information requirements which are subject to review by the Office of Management and Budget (OMB) under section 3504(h) of the Paperwork Reduction Act of 1980. 44 U.S.C. Chapter 35. The Department will submit an information collection request to OMB for its review.

Organizations and individuals desiring to submit comments on this information collection requirement should direct them to the agency official designated for this purpose whose name appears in the preamble and to the Office of Information and Regulatory Affairs, OMB, New Executive Office Building (Room 3208), Washington, DC 20503, Attn.: Desk Officer for HHS.

# List of Subjects in 42 CFR Part 59

Family planning—birth control, Grant programs—health, Health facilities.

Dated: August 28, 1987.

### Robert E. Windom,

Assistant Secretary for Health.

Approved: August 28, 1987.

# Otis R. Bowen,

Secretary.

For the reasons set out in the preamble, it is hereby proposed to amend Subpart A of Part 59, 42 Code of Federal Regulations, as set forth below.

### PART 59-[AMENDED]

 The authority citation for Subpart A of 42 CFR Part 59 is revised to read as follows:

Authority: 42 U.S.C. 300a-4.

2. In 42 CFR 59.2, the following definitions are added:

### § 59.2 [Amended]

"Family planning" means the process of establishing objectives for the number and spacing of a family's children, and selecting the means (including natural family planning methods, adoption, infertility services and general reproductive health care, abstinence and contraception) by which those objectives may be achieved. As such, family planning does not include medical services or counseling related to pregnancy care after pregnancy is diagnosed (including prenatal or postpartum care or counseling), or abortionrelated services. As it relates to the statutory prohibition on the inclusion of abortion as a method of family planning, proper family planning should reduce the incidence of abortion.

"Grantee" means the organization to which a grant is awarded under section 1001 of the Public Health Service Act.

"Organization," as applied to an applicant for or grantee of funds under section 1001 of the Public Health Service Act, means any public or private nonprofit entity in a State. An organization may operate multiple family planning or related programs or projects.

"Program" and "project," which are used interchangeably in these regulations, both refer to the identified activity approved by the Secretary for support under section 1001 of the Public Health Service Act, unless the context indicates otherwise.

"Title X" means Title X of the Public Health Service Act, 42 U.S.C. 300, et seq.

### § 59.5 [Amended]

3. In 42 CFR 59.5, paragraph (a)(5) is removed and paragraphs (a)(6) through

(a)(11) are redesignated as paragraphs (a)(5) through (a)(10) respectively.

# §§ 59.7 through 59.13 [Redesignated as §§ 59.11 through 59.17]

4. In 42 CFR Part 59. §§ 59.7 through 59.13 are redesignated as §§ 59.11 through 59.17 respectively, and new §§ 59.7 through 59.10 are added to read as follows:

# § 59.7 Standards of compliance with prohibition on abortion.

A project may not receive funds under this subpart unless it provides assurance satisfactory to the Secretary that it does not include abortion as a method of family planning. Such assurance must include, at a minmum, representations (supported by documentary evidence. where the Secretary requests) as to compliance with each of the requirements in §§ 59.8 through 59.10. A project supported under this subpart must comply with such requirements at all times during the project period.

# § 59.8 Prohibition on counseling and referral for abortion services; limitation of program services to family planning.

(a) In order to give effect to the statutory prohibition on the use of Title X appropriated funds in projects where abortion is a method of family planning, a project which provides counseling and referral for abortion services as a method of family planning is not eligible to receive funds under this subpart. In addition, because Title X funds are intended only for family planning, services related to pregnancy care after pregnancy is diagnosed may not be provided with Title X funds. Where appropriate, medical or social service referrals for non-Title X supported services shall be made by providing a full list of available health care providers of appropriate prenatal medical care and delivery services and/ or social service agencies from which a family planning client may select. Such referrals may not, however, be used as an indirect means to encourage or promote abortion in violation of section 1008, such as consciously weighting the list of referrals in favor of health care providers and/or facilities which provide abortions. One effect of these regulations will be to insure the ability of otherwise eligible organizations or programs that refuse to engage in abortion-related activities to receive support under this subpart.

(b) Examples. (1) A pregnant client at a family planning clinic supported with Title X funds solicits prenatal care services. Clinic personnel are medically qualified to provide such services. Nonetheless, provision of such services is outside the scope of family planning

supported by Title X.

(2) A client at a family planning clinic supported with Title X funds seeks pregnancy testing and infertility counseling and services. Clinic personnel provide the requested services and in the process thereof discover an ectopic pregnancy. The client is immediately provided a complete list of appropriate hospitals and physicians from which to choose. This service is within the scope of family planning supported by Title X.

(3) A childless husband and wife seek counseling and services relating to infertility and adoption. Such counseling and services are within the scope of family planning supported by Title X.

(4) Clients at a family planning clinic are given a brochure and shown a film about birth control methods that include sections on abortion. Because use of the film and the brochure depicts abortion as a method of family planning, the clinic would not be eligible to receive Title X funds.

# § 59.9 Separation of abortion-related services from family planning programs.

(a) A project supported under this subpart must be kept entirely separate and distinct, financially and physically, from any abortion-related activities. This requirement includes maintaining separate financial, accounting, personnel, and medical record systems and separately maintaining other project functions and physical facilities (including office space, equipment, stationary and the like) in such a manner as to clearly separate Title Xfunded activities from abortion-related activities. This requirement prohibits, by way of example, common waiting, consultation, examination, and treatment areas; shared telephone numbers and receptionists; common names for eligible and ineligible programs; and common office entrances and exits. Although common street or mailing addresses will presumptively constitute a failure to separate adequately Title X-funded programs from other programs which include abortion as a method of family planning, grant applicants may seek to establish the reasonableness of such arrangements in exceptional cases where, as in the example of a large metropolitan hospital with abortion and family planning services located in different wings, the fact of physical separation is otherwise established and no use of appropriated funds in an ineligible program is likely.

(b) Examples. (1) A nonprofit family planning organization operates abortion and family planning clinics

simultaneously on Wednesdays and Fridays in the same one-story building. Nothing on the exterior of the building indicates the existence of two separate programs, although the programs are organized as legally separate entities. The clinics utilize a common parking lot adjacent to the building, a common entrance at the front of the building, and a common receptionist and reception area. The two clinics share the same executive director and financial manager, and the abortion clinic pays a management fee for the services of such personnel. Two other employees of the family planning clinic also work for the abortion clinic. The family planning clinic refers clients to the abortion clinic. The family planning clinic in this example is not "separate and distinct, financially and physically," from abortion-related activities.

(2) A nonprofit organization operates both abortion and family planning clinics at the same address. Both clinics are staffed by the same personnel, and the medical director for the family planning program generally performs the abortions for the abortion clinic as well. The programs, however, schedule clients at different times, with abortion clinic hours only in the mornings and family planning hours only in the afternoon. The schedules do not overlap. The programs use the same telephone number, and the same receptionist answers the phone and makes appointments for both. The programs use the same automobiles, office furnishings, and advertisements. The family planning program in this example is not "separate and distinct, financially and physically," from abortion-related activities.

(3) A private, nonprofit corporation operates a family planning program (Program A) and a program which includes abortion-related services (Program B). Both programs are operated as parts of the same corporate entity, with common directors and officers. Program A and Program B occupy office space leased under the terms of a common master lease, but the offices are in fact located in different sections of the city. Program A maintains entirely separate financial records and has no on-site personnel in common with Program B. The programs conduct no joint advertising and use separate furnishings and equipment. Program A is "separate and distinct, physically and financially," from Program B.

(4) A private, nonprofit organization operates both a family planning clinic that receives Title X funds and an abortion clinic. The clinics are physically separate, but their accounting and financial records are maintained

jointly. Although the family planning clinic is separated physically from the abortion clinic, the joint financial records indicate that the family planning clinic is not "separate and distinct, financially and physically" from abortion-related activities.

(5) A private, nonprofit organization operates both a family planning clinic and an abortion clinic. Both clinics lease space in the same one-story building. The two clinics share a common waiting room. The family planning clinic has separate personnel and maintains separate financial records from the abortion clinic. The family planning clinic in this example is not "separate and distinct, physically and financially" from the abortion-related activities.

# § 59.10 Prohibition on activities that encourage, promote or advocate abortion.

(a) A project supported under this subpart may take no action which encourages, promotes, or advocates abortion as a method of family planning, or which assists a woman in obtaining an abortion as a method of family planning. Actions are considered to encourage, promote, or advocate abortion as a method of family planning if they in any way have the effect of facilitating obtaining abortion as a method of family planning. Such prohibited actions include the following:

(1) Lobbying for the passage of proabortion legislation, providing speakers to argue for abortion as a method of family planning, or paying dues to organizations that advocate abortion as a method of family planning;

(2) Using legal action to make available in any way abortion as a method of family planning:

(3) Developing, assisting in the development of, posting or disseminating in any way materials (including printed matter and audiovisual materials) that advocate abortion as a method of family planning;

(b) Examples. (1) A family planning clinic provides those of its clients who inquire concerning abortion with brochures advertising an abortion clinic. Such a service would "encourage, promote or advocate" abortion.

(2) A family planning clinic pay dues to an organization that devotes a substantial part of its activities to lobbying the Congress for liberalized abortion laws. This activity would "encourage, promote or advocate" abortion.

(3) A family planning clinic displays in its waiting room posters encouraging clients to write their legislative representatives to urge them to vote "pro choice" on pending legislation, and distributes post cards for the same purpose. The clinic is engaged in "encouraging, promoting or advocating" abortion.

(4) A family planning clinic that receives Title X funds assists its clients

in making appointments at abortion clinics. The provision of such services would violate section 1008.

(5) Personnel of a family planning project write their legislative representatives in support of pro-choice legislation, utilizing no project funds to do so. The eligibility of the project for Title X funds would be unaffected by their advocacy of abortion.

[FR Doc. 87-20216 Filed 8-31-87; 8:45 am] BILLING CODE 4160-15-M

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# CFR PARTS AFFECTED DURING SEPTEMBER

At the end of each month, the Office of the Federal Register publishes separately a List of CFR Sections Affected (LSA), which lists parts and sections affected by documents published since the revision date of each title.

# LIST OF PUBLIC LAWS

Note: No public bills which have become law were received by the Office of the Federal Register for inclusion in today's List of Public Laws. Last List August 31,

# TABLE OF EFFECTIVE DATES AND TIME PERIODS—SEPTEMBER 1987

This table is used by the Office of the Federal Register to compute certain dates, such as effective dates and comment deadlines, which appear in agency documents. In computing these

dates, the day after publication is counted as the first day. When a date falls on a weekend or

When a date falls on a weekend or holiday, the next Federal business day is used. (See 1 CFR 18.17) A new table will be published in the first issue of each month.

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September 2	September 17	October 2	October 19	November 2	December 1
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September 14	September 29	October 14	October 29	November 13	December 14
September 15	September 30	October 15	October 30	November 16	December 14
September 16	October 1	October 16	November 2	November 16	December 15
September 17	October 2	October 19	November 2	November 16	December 16
September 18	October 5	October 19	November 2	November 17	December 17
September 21	October 6	October 21	November 5	November 20	December 21
September 22	October 7	October 22	November 6	November 23	December 21
September 23	October 8	October 23	November 9	November 23	December 22
September 24	October 9	October 26	November 9	November 23	December 23
September 25	October 13	October 26	November 9	November 24	December 24
September 28	October 13	October 28	November 12	November 27	December 28
September 29	October 14	October 29	November 13	November 30	December 28
September 30	October 15	October 30	November 16	November 30	December 29